

FORM PTO-1390
(REV 10-94)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

9192.15USWO

U.S. APPLICATION NO. (If known, see 37 C.F.R. 1.5)

Unknown 09/762629

INTERNATIONAL APPLICATION NO.

PCT/IB99/01465

INTERNATIONAL FILING DATE

August 11, 1999

PRIORITY DATE CLAIMED

August 11, 1998

TITLE OF INVENTION

SELECTION METHOD

APPLICANT(S) FOR DO/EO/US

JORSBOE et al.

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(I).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US)
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An unsigned oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment.
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information: 3 Sheets of Formal Drawings; Form PCT/ISA/220; Form PCT/ISA/210; Form PCT/RO/101; Form PCT/IPEA/401; Form PCT/IPEA/101 (Annex); Form PCT/IPEA/408 and attached response; Form PCT/IPEA/416; Form PCT/IPEA/428WP; Form PCT/IPEA/409

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

Unknown **09/762629**

INTERNATIONAL APPLICATION NO

PCT/IB99/01465

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9192.15USWO

17. ☒ The following fees are submitted:**BASIC NATIONAL FEE (37 CFR 1.492(a) (1)-(5)):**

Search Report has been prepared by the EPO or JPO.....\$860.00

International preliminary examination fee paid to USPTO
(37 CFR 1.492(a)(1)).....\$690.00No international preliminary examination fee paid to USPTO (37 CFR 1.482)
but international search fee paid to USPTO (37 CFR 1.445(a)(2)).....\$710.00Neither international preliminary examination fee (37 CFR 1.482) nor
international search fee (37 CFR 1.445(a)(3)) paid to USPTO..... \$1000.00International preliminary examination fee paid to USPTO (37 CFR 1.482)
and all claims satisfied provisions of PCT Article 33(2)-(4)\$100.00**CALCULATIONS PTO USE ONLY****ENTER APPROPRIATE BASIC FEE AMOUNT = \$860.00**Surcharge of **\$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30
months from the earliest claimed priority date (37 CFR 1.492(e)).

\$

CLAIMS

NUMBER FILED

NUMBER EXTRA

RATE

Total claims 48 -20 = 28 X \$18.00 **\$504.00**Independent claims 6 -3 = 3 X \$80.00 **\$240.00**

MULTIPLE DEPENDENT CLAIM(S) (if applicable) + \$260.00 \$

TOTAL OF ABOVE CALCULATIONS = \$1604.00Reduction by 1/2 for filing by small entity, if applicable. Small entity status is claimed
pursuant to 37 CFR 1.27

\$

SUBTOTAL = \$1604.00Processing fee of **\$130.00** for furnishing the English translation later than ☐ 20 ☐ 30
months from the earliest claimed priority date (37 CFR 1.492(f)). + \$**TOTAL NATIONAL FEE = \$1604.00**Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property + \$**TOTAL FEES ENCLOSED = \$1604.00**Amount to be:
refunded \$

charged \$

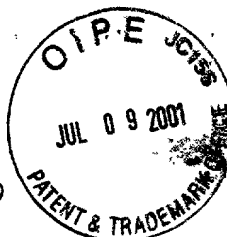
a. ☒ Check(s) in the amount of \$1604.00 to cover the above fees is enclosed.b. ☐ Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any
overpayment to Deposit Account No. 13-2725.**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR
1.137(a) or (b)) must be filed and granted to restore the application to pending status.**

SEND ALL CORRESPONDENCE TO

Dr. Denise Kettelberger
MERCHANT & GOULD
P.O. Box 2903
Minneapolis, MN 55402-0903SIGNATURE: 

NAME: John J. Gresens

REGISTRATION NUMBER: 33,112



PTO/PCT Rec'd 09 JUL 2001

PTO/PCT Rec'd 09 JUN 2001

S/N 09/762,629

PATENT 7

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:	JORSBOE ET AL.	Examiner:	UNKNOWN
Serial No.:	09/762,629	Group Art Unit:	UNKNOWN
Filed:	MAY 14, 2001	Docket No.:	9192.15USWO
Title:	SELECTION METHOD		

CERTIFICATE UNDER 37 CFR 1.8: The undersigned hereby certifies that this correspondence is being deposited in the United States Postal Service, as first class mail, with sufficient postage, in an envelope addressed to: Commissioner for Patents, Washington, D.C. 20231 on July 5, 2001.

By:

Name:

Kristine A. Lueck
Kristine A. Lueck

SUPPLEMENTAL PRELIMINARY AMENDMENT

Box Sequence
Commissioner for Patents
Washington, D.C. 20231

Dear Sir:

Prior to examination please amend the above-identified patent application as follows.

In the Specification

Please replace the paragraph beginning at page 47, line 23, with the following rewritten paragraph:

--In more detail, Figure 3 is the complete nucleotide sequence (SEQ.ID.NO.1) of the *E. coli galT* gene with the deduced amino acid sequence (SEQ.ID.NO.2) shown below the nucleotide sequence. The primers used for the isolation of the gene by PCR are shown with arrows indicating the 5' to 3' direction.--

REMARKS

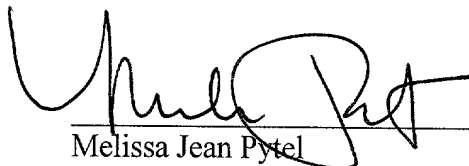
Applicants respectfully request the Examiner to enter the above amendment. No new matter is introduced in the amendment. The changes made to the specification by the current amendment are attached hereto in a page entitled, "Version with Markings to Show Changes Made."

If the Examiner believes a telephone conference would advance the prosecution of this application, the Examiner is invited to telephone the undersigned at the below-listed telephone number.

Respectfully submitted,

MERCHANT & GOULD P.C.
P.O. Box 2903
Minneapolis, Minnesota 55402-0903
(612) 332-5300

Date: July 5, 2001



Melissa Jean Pytel
Reg. No. 41,512
MJP:PSTkaw



VERSION WITH MARKINGS TO SHOW CHANGES MADE

In the Specification

Paragraph beginning at page 47, line 23 has been amended as follows:

In more detail, Figure 3 is the complete nucleotide sequence (SEQ.ID.NO.1) of the *E. coli galT* gene with the deduced amino acid sequence (SEQ.ID.NO.2) shown below the nucleotide sequence. The primers used for the isolation of the gene by PCR are shown with arrows indicating the 5' to 3' direction.

09/762629

JC02 Rec'd PCT/PTO 09 FEB 2001

S/N unknown

PATENTIN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: JORSBOE et al. Serial No.: unknown
 Filed: concurrent herewith Docket No.: 9192.15USWO
 Title: SELECTION METHOD

CERTIFICATE UNDER 37 CFR 1.10

'Express Mail' mailing label number: EL669940715US

Date of Deposit: February 9, 2001

I hereby certify that this correspondence is being deposited with the United States Postal Service 'Express Mail Post Office To Addressee' service under 37 CFR 1.10 on the date indicated above and is addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231.

By: Brian Maharaj
 Name: Brian Maharaj

PRELIMINARY AMENDMENT

Box PCT
 Assistant Commissioner for Patents
 Washington, D. C. 20231

Dear Sir:

In connection with the above-identified application filed herewith, please enter the following preliminary amendments::

IN THE ABSTRACT

Insert the attached Abstract page into the application as the last page thereof.

IN THE SPECIFICATION

A courtesy copy of the present specification is enclosed herewith. However, the World Intellectual Property Office (WIPO) copy should be relied upon if it is already in the U.S. Patent Office.

IN THE CLAIMS

Please cancel claims 1-24 and add new claims 25-73.

T 0 4 T 0 9 " 6 2 9 2 9 4 6 0

25. A method for selecting one or more genetically transformed cells from a population of cells, wherein the population of cells comprising genetically transformed cells comprising a first expressible nucleotide sequence encoding a first expression product, and non-transformed cells wherein the genetically transformed cells are capable of detoxifying a component by action of the first expressible nucleotide sequence or the first expression product; the method comprising:

introducing a medium to the population of cells, the medium comprising a first component or a precursor thereof in an amount that is detoxifiable by the transformed cells but toxic to the non-transformed cells; wherein the first expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and the first component is galactose, a precursor of galactose, or a combination thereof.

26. The method according to claim 25 wherein the genetically transformed cells further comprise a second expressible nucleotide sequence encoding a second expression product wherein the medium comprises a second component or a precursor thereof that is detoxifiable by the genetically transformed cells by the action of the second expressible nucleotide sequence or the second expression product; wherein the second expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose and the second component is galactose, a precursor of galactose, or a combination thereof.

27. The method according to claim 26 wherein the genetically transformed cell further comprises a third expressible nucleotide sequence encoding a third expression product wherein the medium comprises a third component or a precursor thereof that is detoxifiable by the genetically transformed cells by the action of the third expressible nucleotide sequence or the third expression product; wherein the third expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose and the third component is galactose, a precursor of galactose, or a combination thereof.

28. The method according to claim 25 wherein the step of introducing a medium comprising a component to the population of cells comprises introducing a medium comprising a precursor of the component wherein the component is prepared *in situ* from the precursor in the medium.
29. The method according to claim 25 wherein the genetically transformed cells are *in vitro* within a culture.
30. The method according to claim 25 wherein the genetically transformed cells are *in vivo* within an organism.
31. The invention according claim 25 wherein the genetically transformed cells are plant cells.
32. The invention according claim 25 wherein the genetically transformed cells further comprise a nucleotide sequence encoding a nucleotide sequence of interest.
33. A composition comprising a population of cells, the composition comprising:
genetically transformed cells and non-transformed cells wherein the genetically transformed cells comprise a first expressable nucleotide sequence encoding a first expression product; and
a medium comprising a first component or a precursor thereof in an amount that is detoxifiable by the genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product but in an amount that is toxic to the non-transformed cells;
wherein the first expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and wherein the first component is galactose, a derivative of galactose, a precursor of galactose, or a combination thereof.
34. The composition according to claim 33, wherein the genetically transformed cells further comprise a second expressable nucleotide sequence encoding a second expression product,

wherein the medium comprises a second component that is detoxifiable by the genetically transformed cells by action of the second expressible nucleotide sequence or the second expression product; wherein the second expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, a precursor of galactose, or combinations thereof and the second component is galactose, a derivative of galactose, a precursor of galactose, or a combination thereof.

35. The composition according to claim 34, wherein the genetically transformed cells further comprise a third expressible nucleotide sequence encoding a third expression product, wherein the medium comprises a third component that is detoxifiable by the genetically transformed cells by action of the third expressible nucleotide sequence or the third expression product; wherein the third expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, a precursor of galactose, or combinations thereof and the third component is galactose, a derivative of galactose, a precursor of galactose, or a combination thereof.

36. The composition according to claim 33 wherein the medium comprises a precursor of the component and the component is prepared *in situ* from the precursor to provide the component in an amount that is detoxifiable by the transformed cells but toxic to the non-transformed cells.

37. The composition according claim 33 wherein the genetically transformed cells further comprise a nucleotide sequence encoding a nucleotide sequence of interest.

38. The composition according to claim 33 wherein the genetically transformed cells are *in vitro*.

39. The composition according to claim 33 wherein the genetically transformed cells are *in vivo*.

40. The composition according to claim 33 wherein the genetically transformed cells are plant cells.

41. A population of cells comprising genetically transformed cells and non-transformed cells; wherein the genetically transformed cells comprise a first expressable nucleotide sequence encoding a first expression product and are capable of detoxifying a first component by action of the first expressable nucleotide sequence or the first expression product; wherein the first expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and wherein the first component is galactose or a precursor thereof.
42. The population of cells according to claim 41 wherein the genetically transformed cells further comprise a second expressable nucleotide sequence encoding a second expression product and are capable of detoxifying a second component by action of the second expressable nucleotide sequence or the second expression product; wherein the second expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and wherein the second component is galactose or a precursor thereof.
43. The population of cells according to claim 42 wherein the genetically transformed cells further comprise a third expressable nucleotide sequence encoding a third expression product and are capable of detoxifying a third component by action of the third expressable nucleotide sequence or the third expression product; wherein the third expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and wherein the third component is galactose or a precursor thereof.
44. The population of cells according to claim 41 wherein the genetically transformed cells are *in vitro*.
45. The population of cells according to claim 41 wherein the genetically transformed cells are *in vivo*.
46. The population of cells according to claim 41 wherein the genetically transformed cells are plant cells.

47. A genetically transformed cell comprising a first expressable nucleotide sequence encoding a first expression product; wherein the genetically transformed cell is capable of detoxifying a first component by action of the first expressable nucleotide sequence or the first expression product; wherein the first expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose and the first component or precursor thereof is galactose or a precursor thereof.

48. The genetically transformed cell according to claim 47, further comprising a second expressable nucleotide sequence encoding a second expression product wherein the genetically transformed cell is capable of detoxifying a second component by action of the second expressable nucleotide sequence or the second expression product; wherein the second expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and wherein the second component is galactose or a precursor thereof.

49. The genetically transformed cell according to claim 48, further comprising a third expressable nucleotide sequence encoding a third expression product wherein the genetically transformed cell is capable of detoxifying a third component by action of the third expressable nucleotide sequence or the third expression product; wherein the third expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose; and wherein the third component is galactose or a precursor thereof.

50. The genetically transformed cell according to claim 47, wherein the cell is a plant cell.

51. A non-human organism comprising a genetically transformed cell according to claim 47.

52. A plant comprising a genetically transformed cell according to claim 47.

53. The plant according to claim 52, wherein the plant is capable of providing a feed or foodstuff to humans or animals.

54. A plant according to claim 52 wherein the plant is rape seed, potato or maize.
55. A construct for transforming a cell, the construct comprising: a first expressable nucleotide sequence encoding a first expression product, wherein the first expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose.
56. The construct according to claim 55, further comprising a second expressable nucleotide sequence encoding a second expression product, wherein the second expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose.
57. The construct according to claim 56, further comprising a third expressable nucleotide sequence encoding a third expression product, wherein the third expression product is an enzyme capable of metabolizing galactose, a derivative of galactose, or a precursor of galactose.
58. The construct according to claim 55, wherein the cell is a plant cell.
59. A kit for transforming a cell, the kit comprising a construct according to claim 55 and a medium.
60. The kit according to claim 59, wherein the medium comprises a component that is detoxifiable by a transformed cell.
61. A vector comprising the construct according to claim 55.
62. A kit for transforming a cell, the kit comprising a vector according to according to claim 61, and a medium.

transforming some or all of the cells in an initial population of cells with a heterologous nucleotide sequence to form the population of cells containing one or more genetically transformed cells; wherein the heterologous nucleotide sequence is a first expressable nucleotide sequence encoding a first expression product wherein the expression product is an enzyme capable of metabolising galactose, a derivative of galactose, a precursor of galactose; and selecting at least one of the genetically transformed cells using the method of claim 25.

72. The method according to claim 71, wherein the genetically transformed cells further comprise a second expressable nucleotide sequence encoding a second expression product.

73. The method according to claim 72, wherein the genetically transformed cells further comprise a third expressable nucleotide sequence encoding a third expression product.

REMARKS

The above preliminary amendment is made merely to conform the claim structure to U. S. practice. A new abstract page is supplied to conform to that appearing on the publication page of the WIPO application, but the new Abstract is typed on a separate page as required by U.S. practice.

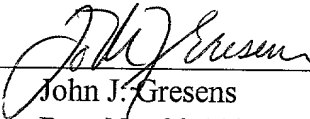
Applicants respectfully request that the preliminary amendment described herein be entered into the record prior to calculation of the filing fee and prior to examination and consideration of the above-identified application.

If a telephone conference would be helpful in resolving any issues concerning this communication, please contact Applicants' primary attorney-of record, Dr. Denise Kettelberger (Reg. No. 33,924), at 612.371.5268.

Respectfully submitted,

MERCHANT & GOULD P.C.
P.O. Box 2903
Minneapolis, Minnesota 55402-0903
(612) 332-5300

Dated: February 9, 2001

By 
John J. Gresens
Reg. No. 33,112

JJG/kas

097629.05404
T 04 T 50 6 29 29 260

A selection method for selecting from a population of cells one or more selectable genetically transformed cells is described. The population of cells comprises selectable genetically transformed cells and possible non-transformed cells. Each of the selectable

10 genetically

transformed cells comprises a first expressable nucleotide sequence encoding a first expression product; and optionally a second expressable

nucleotide sequence encoding a second expression product and/or a

third expressable nucleotide sequence encoding a third expression product. A component

15 is utilisable by the selectable genetically transformed cells by action of the first

expressable nucleotide sequence or the first expression product and optionally by action of the optional

second expressable nucleotide sequence or the optional second expression

product and/or by, action of the optional third expressable nucleotide sequence or the

20 optional Third expression product. The component can

be present in an amount that is toxic to the non-transformed cells.

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SELECTION METHOD

The present invention relates to a selection method.

- 5 The present invention also relates to an enzyme and a nucleotide sequence coding for same that are useful in a selection method.

In particular, the present invention relates to a method for the selection (e.g. identification and/or separation) of genetically transformed cells and compounds and
10 genetic material for use in the method.

It is well known that when a nucleotide sequence of interest ("NOI") is to be introduced into a population of cells by transformation, only a certain number of the cells are successfully transformed, i.e., only a certain number of the cells receive the NOI. It is
15 then necessary to identify the genetically transformed cells so that these cells may be separated from the non-transformed cells in the population. For the production of transgenic plants etc., this often requires the use of a selection system that allows the regeneration and growth of the transformed (or transgenic) cells. As these transformed cells frequently constitute a minor fraction of the treated cells, compared to the majority
20 of cells which remain untransformed, so the selection system has to be able to be effective in selecting out the transformed cells.

A common technique for a selection method includes introducing transformed cells and non-transformed cells into a medium that comprises a substance which the transformed
25 cells are able to tolerate. In that medium the transformed cells are able to survive and grow, while the non-transformed cells are prone to growth inhibition and, in some cases, are killed.

Thus, to date, the general strategy has been to introduce a selectable gene along with the
30 NOI(s), and then allowing the transformed cells to survive on selective media while the non-transformed cells are killed (Bowen 1993).

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Typically, if a population of plant cells has been subjected to genetic transformation, selection of the transformed cells typically takes place using a selection gene which codes for antibiotic resistance or herbicide resistance. The selection gene is coupled to or co-introduced with the NOI to be incorporated into the plant in question, so that both of the two sequences are incorporated into some or all of the population of cells.

As not all of the cells may have been transformed, the cells are then cultivated on or in a medium containing the respective antibiotic or herbicide to which the genetically transformed cells are resistant by virtue of the selection gene. In this medium, the transformed cells are able to grow and thus be identified out of the total cell population, since the non-transformed cells - which do not contain the antibiotic or herbicide resistance gene in question - have an inhibited growth or even are killed.

So far, the most widely used selectable gene is the neomycin phosphotransferase II (NPTII) gene (Fraley *et al.* 1986) which confers resistance to the aminoglycoside antibiotics kanamycin, neomycin and G-418 (Bevan *et al.* 1983). A number of other selective systems has been developed based on resistance to bleomycin (Hille *et al.* 1986), bromoxynil (Stalker *et al.* 1988), chloramphenicol (Fraley *et al.* 1983), 2,4-dichlorophenoxy-acetic acid (Streber and Willmitzer 1989), glyphosate (Shah *et al.* 1986), hygromycin (Waldron *et al.* 1985) or phosphinothricin (De Block *et al.* 1987).

The selection methods which rely on the use of antibiotics or herbicides suffer from a number of disadvantages. For example, there is concern amongst some people, such as environmental groups and governmental authorities, as to whether it is environmentally safe to incorporate genes coding for antibiotic resistance and/or herbicide resistance into plants and micro-organisms. This concern is of particular significance for food plants and for micro-organisms which are not designed and/or intended to be used in a closed environment (e.g. micro-organisms for use in agriculture), and also for micro-organisms which are designed for use in a closed environment but which may be released from the closed environment.

While such ecological concerns may prove unfounded, as suggested by Flavell *et al.* (1992), they may nevertheless lead to governmental restrictions on the use of antibiotic resistance genes in transgenic plants, and it is therefore desirable to develop new selection methods which are independent of such genes.

In addition, in some or many cases, the corresponding antibiotic or herbicide resistance genes may not be relevant to the desired transgenic trait. Also, they may be undesirable in the final product (Yoder and Goldsbrough 1994).

Thus, the use and subsequent release of selectable genes such as antibiotic resistance genes into the environment has been the target of concern among environmental authorities.

Therefore, new selection systems for the production of transgenic plants without any herbicide or antibiotic resistance genes have been developed. By way of example, these new selection systems include three selection systems based on the concept of 'positive selection' wherein transgenic cells acquire a gene which confers a metabolic advantage to those cells whilst non-transgenic cells starved rather than killed. Some of these selection systems have been reviewed by Joersbo (Joersbo 1997).

One of these positive selection methods relates to cytokinins which must be added to obtain optimal shoot regeneration. By adding cytokinin as an inactive glucuronide derivative and using a β -glucuronidase gene as selectable gene, cells which have acquired this gene by transformation are able to convert the cytokinin glucuronide to active cytokinin while untransformed cells are arrested in development (Joersbo and Okkels 1996; Okkels *et al.* 1997).

Two other selection systems employ, as selectable agents, the carbohydrates mannose and xylose, which are not metabolised by a number of plant species (Bojsen *et al.* 1994). By substituting the normally employed carbohydrate with one of these compounds, cells transformed with a gene encoding an enzyme capable of converting it to a metabolisable

isomer are favoured in growth while the non-transgenic cells are starved. Mannose is initially phosphorylated to mannose-6-phosphate by hexokinase present in plant cells but this compound is not metabolised any further in many plant species. Cells transformed with a phosphomannose isomerase gene are able to convert mannose-6-phosphate to the readily metabolised fructose-6-phosphate, giving these cells a metabolic advantage (Joersbo *et al.* 1998). Xylose can be converted to xylulose by xylose isomerase which functions as the selectable marker in this system (Haldrup 1996).

Another selection system that is not dependent on the use of herbicide and antibiotic resistance genes is described in PCT/GB98/00367. In the general method of PCT/GB98/00367, selectable transformed cells are selected from a population of cells which comprises the selectable genetically transformed cells and possible non-transformed cells. Each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence and optionally a second expressable nucleotide sequence. In the selection method, a component or a metabolic derivative thereof when present in a low concentration in a medium is a nutrient for both the selectable genetically transformed cells and the non-transformed cells. In the selection method, the component or the metabolic derivative thereof when present in a high concentration in a medium is toxic to the non-transformed cells. The first nucleotide sequence codes for a gene product capable of converting the component or the metabolic derivative thereof when present in a high concentration in a medium to a nutrient for the selectable genetically transformed cells. The selection method comprises the step of introducing the population of cells to a medium, wherein the medium optionally comprises a high concentration of the component or the metabolic derivative thereof. In the selection method, the component or the metabolic derivative thereof is a source of both carbohydrate and nitrogen for the selectable genetically transformed cells. Alternatively, in the selection method if a portion of the component serves as a metabolic substrate and is metabolically converted to a derivatised substrate, then that derivatised substrate is capable of providing an allosteric effect on the gene product. In one preferred aspect, the selection method relies on the use of glucosamine.

Despite the advent of selection methods that do not necessarily rely on the use of antibiotic or herbicide resistance genes, it is still desirable to develop new methods for selecting genetically transformed cells or organisms (or parts thereof) comprising such.

5 According to a first aspect of the present invention there is provided:

a selection method for selecting from a population of cells one or more selectable genetically transformed cells,

wherein the population of cells comprises selectable genetically transformed cells and possible non-transformed cells;

10 wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third
15 expression product;

wherein a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable
20 nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

the method comprising the step of introducing the population of cells to a medium,

wherein the medium comprises the component and/or a derivative thereof and/or
25 a precursor thereof and in an amount such that the component is or will be in an amount that is utilisable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an
30 enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

For convenience, we sometimes refer to the selection method of the present invention as being the "galactose selection method". However, it is to be understood that the method of the present invention is not necessarily limited to galactose as being the component or the precursor of the component that enables workers to select transformed cells over non-transformed cells. However, and as will become apparent, in a preferred embodiment galactose is a preferred component or precursor for the component and/or galactose-1-phosphate is a preferred component or derivative thereof.

The component may be any suitable chemical compound, product, molecule etc.

The term "utilisable" as used herein with reference to the present invention (e.g. "the component is utilisable by the selectable genetically transformed cells") means that the component can be processed in a non-adverse or non-detrimental fashion by the selectable genetically transformed cells. Thus, the term includes metabolising (such as metabolising in a beneficial manner), detoxifying, etc. For some applications, the term "utilisable" may mean at least "metabolisable". For some applications, the term "utilisable" may mean at least "detoxifiable". Here, the term "detoxifiable" means capable of being converted to one or more derivatives that do not have an adverse effect on the transformed cell. An example of an adverse effect is growth inhibition. These derivatives may or may not accumulate in the cells. In addition, or in the alternative, these derivatives may or may not be fully metabolised by the transformed cell.

The derivative of the component can be any suitable derivative - such as a chemical derivative, a metabolic derivative, etc.

For some applications, preferably the derivative of the component is at least a metabolic derivative.

The precursor of the component can be any suitable precursor - such as a metabolic precursor, etc.

For some applications, preferably the precursor of the component is at least a metabolic precursor.

The action of the the first expressable nucleotide sequence or the first expression product and optionally the optional second expressable nucleotide sequence or the optional second expression product and/or the optional third expressable nucleotide sequence or the optional third expression product may be direct or indirect. Here, indirect action may mean that the first expression product or the optional second expression product or the optional third expression product converts a precursor of the component to, for example, the component or even another precursor thereof or a derivative thereof. However, the component that is present or is produced by direct or indirect action is or will be in an amount that is utilisable by the transformed cells but in an amount that is toxic to the non-transformed cells.

According to a second aspect of the present invention there is provided

a composition comprising a population of cells comprising selectable genetically transformed cells and possible non-transformed cells; and a medium;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

wherein the medium comprises the component and/or a derivative thereof and/or a precursor thereof and in an amount such that the component is or will be in an amount that is utilisable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

According to a third aspect of the present invention there is provided

a population of cells comprising selectable genetically transformed cells and possible non-transformed cells;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an

enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof;
and

wherein the component and/or the derivative thereof and/or the precursor thereof
is galactose or a derivative thereof or a precursor thereof.

5

According to a fourth aspect of the present invention there is provided

a selectable genetically transformed cell comprising a first expressible nucleotide
sequence encoding a first expression product; and optionally a second expressible
nucleotide sequence encoding a second expression product and/or a third expressible
nucleotide sequence encoding a third expression product;

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wherein a component is utilisable by the selectable genetically transformed cells
by action of the first expressible nucleotide sequence or the first expression product and
optionally by action of the optional second expressible nucleotide sequence or the
optional second expression product and/or by action of the optional third expressible
nucleotide sequence or the optional third expression product;

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wherein the component can be present in an amount that is toxic to the non-
transformed cells;

wherein each of the first expression product and the optional second expression
product and the optional third expression product is independently selected from an
enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof;
and

20

wherein the component and/or the derivative thereof and/or the precursor thereof
is galactose or a derivative thereof or a precursor thereof.

25

According to a fifth aspect of the present invention there is provided an organism
comprising a selectable genetically transformed cell according to the present invention.

According to a sixth aspect of the present invention there is provided

a construct for genetically transforming a non-transformed cell to produce a
selectable genetically transformed cell;

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the construct comprising a first expressable nucleotide sequence encoding a first expression product; and optionally a second expressable nucleotide sequence encoding a second expression product and/or a third expressable nucleotide sequence encoding a third expression product;

5 wherein a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

10 wherein the component can be present in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof;

15 and

wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

According to a seventh aspect of the present invention there is provided a vector
20 comprising the construct according to the present invention.

According to an eighth aspect of the present invention there is provided a plasmid comprising the construct according to the present invention.

25 According to a ninth aspect of the present invention there is provided a kit comprising a construct according to the present invention or a vector according to the present invention or a plasmid according to the present invention for genetically transforming a non-transformed cell to produce a selectable genetically transformed cell; and a medium.

According to a tenth aspect of the present invention there is provided a plant or plant cell comprising one or more heterologous enzymes, wherein the heterologous enzymes are any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2).

Here the term "heterologous enzymes" includes enzymes that are derived from a different species or even a different cell. The term also includes enzymes (which may or may not be native to the cell) which are expressed by recombinant nucleotide sequences.

The term also includes homologous enzymes that have been expressed by homologous coding sequences but when under the control of heterologous promoters. Here the term "heterologous promoters" means promoters that are not naturally associated with the coding sequence in question.

According to an eleventh aspect of the present invention there is provided a feed, foodstuff or food prepared from or comprising the aspects of the present invention.

According to a twelfth aspect of the present invention there is provided the use of any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) as a selection means for selecting a genetically transformed cell over a non-transformed cell.

According to a thirteenth aspect of the present invention there is provided the use of any one or more of a nucleotide sequence coding for galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) for providing a selection means for selecting a genetically transformed cell over a non-transformed cell.

According to a fourteenth aspect of the present invention there is provided the use of any one or more of galactose, galactose-1-phosphate, UDP-galactose, or derivatives thereof as a selection means for selecting a genetically transformed cell over a non-transformed cell.

According to a fifteenth aspect of the present invention there is provided

a selection method for selecting from a population of cells one or more selectable genetically transformed cells,

wherein the population of cells comprises selectable genetically transformed cells and possible non-transformed cells;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

the method comprising the step of preparing the population of cells by transforming some or all of the cells in an initial population of cells containing non-transformed cells with a heterologous nucleotide sequence so as to form the population of cells containing one or more selectable genetically transformed cells; and

selecting at least one of the selectable genetically transformed cells;

wherein the heterologous nucleotide sequence is any one or more of the first expressable nucleotide sequence, the optional second expressable nucleotide sequence or the optional third expressable nucleotide sequence;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof.

Here, the term "heterologous nucleotide sequence" includes nucleotide sequences that are derived from a different species or even a different cell. The term also includes nucleotide sequences (which may or may not be native to the cell) that have been

prepared by use of recombinant DNA techniques. The term also includes native nucleotide sequences but when under the control of heterologous promoters. Here the term "heterologous promoters" means promoters that are not naturally associated with the nucleotide sequence in question.

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In one preferred aspect, the selectable genetically transformed cell/cells is/are *in vitro* within a culture.

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In an alternative preferred aspect, the selectable genetically transformed cell/cells is/are *in vivo* within an organism.

Preferably the selectable genetically transformed cell/cells is/are selectable genetically transformed plant cell/cells.

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Preferably an additional nucleotide sequence is present and wherein the additional nucleotide sequence codes for a nucleotide sequence of interest ("NOI").

Preferably the organism is a plant.

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Preferably the plant is capable of providing a foodstuff for humans or animals.

In an alternative preferred aspect, the plant is capable of providing a commodity for humans - such as cotton, tobacco etc.

25

Preferably the plant (or part thereof, including cells thereof) is a monocot or a dicot (including legumes).

In a preferred aspect, the plant is any one of rape seed, potato or maize.

30

For some applications, preferably the component is present in the medium.

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For some applications, preferably the component is prepared *in situ* in the cell from a precursor that was present in the medium.

If the component is prepared *in situ* in the cell from a precursor that was present in the medium, then galactose is a preferred component or precursor for the component and/or a preferred component or derivative thereof is galactose-1-phosphate.

More preferably, if the component is prepared *in situ* in the cell from a precursor that was present in the medium, then galactose is a preferred precursor for the component and/or a preferred component is galactose-1-phosphate.

Thus, in each aspect of the present invention, a precursor for the component is preferably metabolically converted to the component by the transformed cell.

When the component or the derivative thereof or precursor thereof is present in the medium then preferably the component or the derivative thereof or precursor thereof is present in an amount that does not detrimentally affect a major proportion of the transformed cells.

Preferably, when the component or the derivative thereof or precursor thereof is present in the medium then the component or the derivative thereof or precursor thereof is present in an amount that does not detrimentally affect substantially most of the transformed cells.

More preferably, when the component or the derivative thereof or precursor thereof is present in the medium then the component or the derivative thereof or precursor thereof is present in an amount that does not detrimentally affect substantially all of the transformed cells.

In a further aspect, in some cases the medium need not contain any added quantities of the component or the derivative or precursor thereof according to the present invention.

Preferably, however, the medium contains added quantities of the component or the derivative or precursor thereof according to the present invention.

- 5 For some applications, if carbohydrates are also present in addition to the component or the derivative thereof or the precursor therefor then preferably the carbohydrates are present in amounts that do not affect substantially the effect of the component or the derivative thereof or the precursor therefor. Typically, said levels of carbohydrate(s) will be low levels.

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Other aspects of the present invention include:

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The use of any one or more of galactose, galactose-1-phosphate, UDP-galactose, or any derivative thereof, as a selection means for selecting a genetically transformed cell over a non-transformed cell.

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The use of any one or more of a nucleotide sequence coding for any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) for providing a selection means for selecting a genetically transformed cell over a non-transformed cell.

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The use of any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) for providing a selection means for selecting a genetically transformed cell over a non-transformed cell.

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In the above aspects, the phrase "selecting a genetically transformed cell over a non-transformed cell" can be alternatively expressed as "selecting a genetically transformed cell from one or more non-transformed cells".

Preferably, any one or more of the first, second or third expressable nucleotide sequence and/or any one or more of the first, second or third expression product (such as any one or more of the enzymes galactokinase, UTP-dependent pyrophosphorylase, UDP-glucose dependent uridylyltransferase and UDP-galactose epimerase and/or nucleotide sequences coding for same) are, for example, obtainable from any one or more of: *E. coli* (CE Vorgias, H-G Lemaire, KS Wilson 1991. Protein Expr. and Pur. 2, 330-338); *Saccharomyces sp.* (RA Darrow, R Rodstrom 1968. Biochemistry 7, 1645-1654; MA Schell, DB Wilson 1977. J. Biol. Chem. 252, 1162-1166); *Streptomyces coelicolor* A3 (BD Wilson, DS Hogness 1966. in Methods in Enzymol. vol. 8, pp. 229-240, Academic Press, San Diego); *Tetrahymena thermophila* (JE Lavine, E Cantlay, CT Roberts, DE Morse 1982. Biochim. Biophys. Acta 717, 76-85); *Clostridium pasteurianum* (F Daldal, J Applebaum 1985. J. Mol. Biol. 186, 533-545); *Kluveromyces lactis* (MI Riley, RC Dickson 1984. J. Bacteriol. 158, 705-712); *Vicia faba*; *Petunia sp.*; and mammals eg. Chinese hamster (B Talbot, J-P Thirion 1982. Int. J. Biochem. 14, 719-725), human (WO 96/09374).

According to one aspect of the present invention there is provided a selection system that uses a component or a precursor therefor or a derivative thereof for selecting at least one genetically transformed cell from a population of cells, wherein the at least one genetically transformed cell is transformed with a nucleotide sequence which encodes an expression product capable of utilising the component but wherein the component toxic to non-transformed cells.

According to another aspect of the present invention there is provided a selection system for selecting at least one genetically transformed cell from a population of cells in a medium, wherein the at least one genetically transformed cell is transformed with a nucleotide sequence which encodes an expression product capable of converting a component or precursor therefor or a derivative thereof, that is present in the medium that provides the component at an amount that is toxic to non-transformed cells but is utilisable (e.g. metabolisable and/or detoxifiable) by the at least one transformed cell.

The term "cells" is intended to refer to any type of cells from which individual genetically transformed cells may be identified and isolated using the method of the invention. Examples of such cells typically include cells of plants that have a commercial worth - such as crops useful for food or feed production. If desired, other plant cells can be transformed. The term "cells" is also meant to encompass protoplasts, i.e. the protoplasm of a cell enclosed in a membrane but without a cell wall. While it is contemplated that the selection method of the present invention may be used for any type of cell, the method has been found to be particularly suitable for the selection of genetically transformed plant cells.

10

The term "population of cells" refers to any group of cells which has been subjected to genetic transformation and from which it is desired to identify those cells which have been genetically transformed and to isolate the genetically transformed cells from non-genetically transformed cells. The population may, for example, be a tissue, an organ or a portion thereof, a population of individual cells in or on a substrate, such as a culture of plant cells, for example a population of cells in a solution or suspension, or a whole organism, such as an entire plant.

15

The term "selecting" refers to the process of identifying and/or isolating the genetically transformed cells from the non-genetically transformed cells using the method of the present invention.

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The term "medium" includes any medium that is capable of sustaining the viability (such as growth) of the transformed cells while inhibiting or killing the non-transformed cells. For example, the medium may comprise typical ingredients of a growth medium but wherein those ingredients are in such an amount that the transformed cells are selectively grown. In accordance with the present invention, the medium may typically comprise at least a component, or a precursor therefor or a derivative thereof, according to the present invention.

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The term "toxic" as used herein in relation to the non-transformed cells means that the component or derivative thereof or a precursor thereof has an adverse effect on the non-transformed cells or is metabolised to a derivative that has an adverse effect on the non-transformed cells. An example of an adverse effect is growth inhibition. The term also includes death of the non-transformed cells.

The term "genetically transformed" includes transformation using recombinant DNA techniques.

The term "introducing the population of cells to a medium" means adding the population of cells to the medium or *vice versa*.

The component of the present invention may be derived from a precursor therefor.

Preferably, the precursor is present in a medium and the precursor is processed by the cells to the component - such as by use of naturally occurring enzymes and/or by use of heterologous enzymes (such as those expressed by recombinant DNA).

The term "vector" includes expression vectors and transformation vectors.

The term "expression vector" means a construct capable of *in vivo* or *in vitro* expression.

The term "transformation vector" means a construct capable of being transferred into a species - such as a plant by use of, for example, direct gene transfer techniques (such as particle gun techniques for monocot transformation).

The term "tissue" includes tissue *per se* and organ.

The term "organism" in relation to the present invention includes any organism that could comprise the nucleotide sequence coding for the enzyme according to the present invention and/or products obtained therefrom, and/or wherein the nucleotide sequence according to the present invention can be expressed when present in the organism.

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Preferably the organism is a plant.

The term "transgenic organism" in relation to the present invention includes any organism that comprises the nucleotide sequence coding for the enzyme according to the present invention and/or products obtained therefrom, and/or wherein the nucleotide sequence according to the present invention can be expressed within the organism. Preferably the nucleotide sequence is incorporated in the genome of the organism.

Preferably the transgenic organism is a plant.

15

In a highly preferred embodiment, the transgenic organism (or part thereof) does not comprise the combination of a promoter and at least one of the first nucleotide sequence coding for the first enzyme according to the present invention, the second nucleotide sequence coding for the second enzyme according to the present invention and the third nucleotide sequence coding for the third enzyme according to the present invention, wherein both the promoter and the nucleotide sequence are native to that organism (or part thereof) and are in their natural environment. Thus, in this highly preferred embodiment, the present invention does not cover at least one of a native first nucleotide sequence coding for the first enzyme according to the present invention, a native second nucleotide sequence coding for the second enzyme according to the present invention and a native nucleotide sequence coding for the third enzyme according to the present invention in its natural environment when it is under the control of its native promoter which is also in its natural environment. In addition, in this highly preferred embodiment, the present invention does not cover the native enzyme according to the present invention (namely any one of the first enzyme according to the present invention, the second enzyme according to the present invention, the third enzyme according to the

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the present invention means at least not causing an adverse, detrimental effect to the transformed cells.

In accordance with the present invention, a non-transformed cell is a cell that does not
5 comprise at least one of the first nucleotide sequence according to the present invention, the second nucleotide sequence according to the present invention and the third nucleotide sequence according to the present invention.

The non-transformed cell of the present invention may even be a previously transformed
10 cell that does not comprise at least any one of the first nucleotide sequence according to the present invention, the second nucleotide sequence coding according to the present invention and the third nucleotide sequence according to the present invention. The non-transformed cell of the present invention can, however, contain one or more heterologous nucleotide sequences, or one or more homologous nucleotide sequences
15 under the control of one or more heterologous transcriptional control elements.

The transformed cell of the present invention may contain one or more heterologous nucleotide sequences, or one or more homologous nucleotide sequences under the control of one or more heterologous transcriptional control elements.

In a highly preferred embodiment, the first nucleotide sequence is not in its natural environment. In this regard, the first nucleotide sequence may not be native (i.e. foreign) to the cell or organism. In addition, the first nucleotide sequence may be native to the cell or organism but wherein the first nucleotide sequence is operably linked to a
20 promoter that is heterologous to the first nucleotide sequence.
25

In accordance with the present invention there may be a plurality of first nucleotide sequences and/or second nucleotide sequences and/or third nucleotide sequences.

The or each first nucleotide sequence may be independently selected from DNA or RNA. Preferably, the or each first nucleotide sequence is DNA. More preferably, the or each first nucleotide sequence is recombinant DNA.

- 5 The second nucleotide sequence may be independently selected from DNA or RNA. Preferably, the or each second nucleotide sequence is DNA. More preferably, the or each second nucleotide sequence is recombinant DNA.

- The third nucleotide sequence may be independently selected from DNA or RNA.
10 Preferably, the or each third nucleotide sequence is DNA. More preferably, the or each third nucleotide sequence is recombinant DNA.

- The term "recombinant DNA" means DNA prepared by at least one step that utilises at least one recombinant DNA technique.

- 15 Thus, in a first broad aspect the present invention relates *inter alia* to the use of at least one nucleotide sequence encoding an expression product that affects galactose or a derivative thereof or a precursor thereof as a selective agent for the selection of transgenic cells, tissues, organs, organisms (such as plants).

- 20 In a second broad aspect the present invention relates *inter alia* to the use of the expression product of at least one nucleotide sequence that affects galactose or a derivative thereof or a precursor thereof as a selective agent for the selection of transgenic cells, tissues, organs, organisms (such as plants).

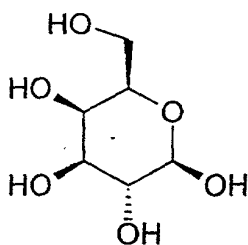
- 25 In a third broad aspect the present invention relates *inter alia* to galactose or a derivative thereof or a precursor thereof as a selective agent for the selection of transgenic cells, tissues, organs, organisms (such as plants).

- 30 Galactose is a hexose which has been demonstrated to be toxic to most plant species (eg. Farkas 1954; Hughes and Street 1974; Roberts et al. 1971).

Galactose has optical isomeric forms and can occur as a linear or a cyclic structure.

An example of a galactose molecule can be pictorially represented as:

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Preferably the component is galactose, preferably D-galactose.

- 10 If the medium of the present invention comprises a precursor of galactose then preferably the precursor is capable of liberating free galactose or a derivative thereof.

Preferably the precursor is capable of liberating free galactose.

- 15 More preferably the precursor is capable of liberating free D-galactose.

If the medium of the present invention comprises a precursor of galactose then preferably one or more enzymes are present which can liberate free galactose or a derivative thereof from the precursor.

20

Preferably if the medium of the present invention comprises a precursor of galactose then preferably one or more enzymes are present which can liberate free D-galactose.

Preferably the enzyme(s) is at least a galactosidase.

25

Preferably the enzyme(s) is/are selected from one or more of α -galactosidases or β -galactosidases.

The enzyme can be present in the medium already or it can be prepared by the medium itself. Alternatively, the enzyme can be prepared by the transformed cell. Here, the enzyme can be native to the cell or it can be heterologous to the cell. The transformed cell may also comprise an additional nucleotide sequence coding for an enzyme capable of releasing galactose from a precursor thereof.

Preferable examples of precursors of galactose include galactose containing compounds - such as lactoses, melibioses, raffinoses, stachyoses, verbascoses and galactinols.

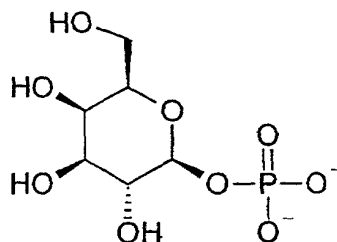
More preferable precursors of galactose include α -lactose (β -D-galactopyranosyl [1 \rightarrow 4]- α -D-glucose, milk sugar), β -lactose, melibiose (6-O- α -D-galacto-pyranosyl-D-glucose), raffinose, stachyose, verbascose and galactinol and any other substrate which liberates free D-galactose upon hydrolysis by either α -galactosidases or β -galactosidases.

Other examples of potentially useful precursors for use in the galactose selection method of the present invention are chemically derivatised forms of galactose, preferably chemical derivatives of D-galactose, from which free galactose can be liberated by use of appropriate techniques, such as enzyme action. By way of example, suitable chemical derivatives are D-galactose pentaacetate and D-galactose methyl galactoside.

Alternatively or in addition, the medium may comprise a derivative of galactose.

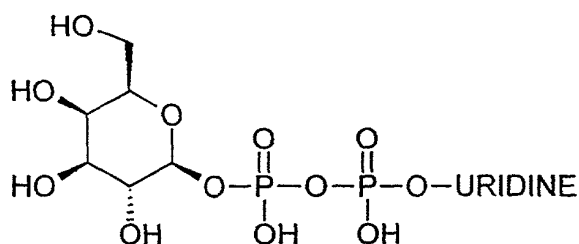
Preferable examples of such a derivative include galactose-1-phosphate and UDP-galactose.

Galactose-1-phosphate can be pictorially represented as:



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UDP-galactose can be pictorially represented as:



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More preferable examples of such a derivative include D-galactose-1-phosphate and UDP-D-galactose.

15

Preferably the component is galactose-1-phosphate.

Preferably the precursor for the component is galactose.

- 20 Preferably each of the first expression product; the second expression product and the third expression product is independently selected from galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2).

Preferably each of the first expression product; the second expression product and the third expression product is independently selected from galactokinase (EC 2.7.1.6) obtainable from Petunia, yeast and other microorganisms, *Vicia faba*, *Phaseolus aureus*,
5 barley or corn, UTP-dependent pyrophosphorylase (EC 2.7.7.10) obtainable from Petunia, UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12) obtainable from yeast and other microorganisms, UDP-galactose epimerase (EC 5.1.3.2) obtainable from Petunia, yeast and other microorganisms, fenugreek, wheat, sugarcane and various trees.

10 Preferably, each of the first expressable nucleotide sequence, the second expressable nucleotide sequence and the third expressable nucleotide is independently selected from expressable nucleotide sequences coding for any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2).

15 Preferably, each of the first expressable nucleotide sequence, the second expressable nucleotide sequence and the third expressable nucleotide is independently selected from expressable nucleotide sequences coding for any one or more of galactokinase (EC 2.7.1.6) obtainable from Petunia, yeast and other microorganisms, *Vicia faba*, *Phaseolus*
20 *aureus*, barley or corn, UTP-dependent pyrophosphorylase (EC 2.7.7.10) obtainable from Petunia, UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12) obtainable from yeast and other microorganisms, UDP-galactose epimerase (EC 5.1.3.2) obtainable from Petunia, yeast and other microorganisms, fenugreek, wheat, sugar cane and various trees.

25 In one preferred aspect of the present invention, the transformed cells comprise the first expressable nucleotide sequence but not the optional second expressable nucleotide sequence and not the optional third expressable nucleotide sequence.

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In this aspect of the present invention, preferably the first expressable nucleotide sequence is a nucleotide sequence coding for UDP-galactose epimerase (EC 5.1.3.2) or UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12). In this aspect, preferably the medium comprises UDP-galactose and/or galactose-1-phosphate, or derivatives thereof.

5

In one preferred aspect of the present invention, the transformed cells comprise the first expressable nucleotide sequence and the optional second expressable nucleotide sequence, but not the optional third expressable nucleotide sequence.

10 In this aspect of the present invention, preferably the first expressable nucleotide sequence is a nucleotide sequence that codes for UDP-galactose epimerase, and the second expressable nucleotide sequence is a nucleotide sequence that codes for UTP-dependent pyrophosphorylase (EC 2.7.7.10) and/or UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12). In this aspect, preferably the medium comprises at
15 least galactose-1-phosphate. For example, the medium may comprise galactose-1-phosphate and UDP-galactose, or derivatives thereof.

In one preferred aspect of the present invention, the transformed cells comprise the first expressable nucleotide sequence and the optional third expressable nucleotide sequence,
20 but not the optional second expressable nucleotide sequence.

In this aspect of the present invention, preferably the first expressable nucleotide sequence is a nucleotide sequence that codes for UDP-galactose epimerase, and the third expressable nucleotide sequence is a nucleotide sequence that codes for galactokinase
25 (EC 2.7.1.6). In this aspect, preferably the medium comprises at least galactose. For example, the medium may comprise galactose and UDP-galactose, or derivatives thereof.

In one preferred aspect of the present invention, the transformed cells comprise the first
30 expressable nucleotide sequence, the optional second expressable nucleotide sequence, and the optional third expressable nucleotide sequence.

In this aspect of the present invention, preferably the first expressable nucleotide sequence is a nucleotide sequence that codes for UDP-galactose epimerase, the second expressable nucleotide sequence is a nucleotide sequence that codes for UTP-dependent pyrophosphorylase (EC 2.7.7.10) and/or UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), and the third expressable nucleotide sequence is a nucleotide sequence that codes for galactokinase (EC 2.7.1.6). In this aspect, preferably the medium comprises at least galactose, or derivatives thereof. For example, the medium may comprise galactose and galactose-1-phosphate, or derivatives thereof. For example, the medium may comprise galactose and UDP-galactose, or derivatives thereof. For example, the medium may comprise galactose, galactose-1-phosphate and UDP-galactose, or derivatives thereof.

In accordance with the present invention, the medium can comprise one or more of the component, the derivative thereof and the precursor.

In order to prepare any one or more of the transformed cells, tissues, organs and organisms according to the present invention, the first expressable nucleotide sequence, the optional second expressable nucleotide sequence and the optional third expressable nucleotide sequence can be introduced into the original non-transformed cells by use of any one or more of a single construct, a single plasmid, or a single vector. Alternatively, the first expressable nucleotide sequence, the optional second expressable nucleotide sequence and the optional third expressable nucleotide sequence can be introduced into the original non-transformed cells by use of any one or more of: two or more constructs, two or more plasmids, or two or more vectors. --

In accordance with the present invention any one or more of the first nucleotide sequence, the second nucleotide sequence and the third nucleotide sequence may comprise an intron.

In this respect, the presence of the intron within the coding portions of the first nucleotide sequence and/or the second nucleotide sequence and/or the third nucleotide sequence may inactivate the expression products thereof *vis-a-vis* a prokaryote, such as a prokaryotic vector system used to generate the transformed cells, tissue, organs or organism.

The present invention therefore provides a method for selecting genetically transformed cells - such as cells into which a NOI has been incorporated - by providing the transformed cells with a selective advantage.

The method of the present invention is not dependent on the preparation of genetically transformed plants containing as a selection means a nucleotide sequence coding for antibiotic or herbicide resistance. Nevertheless, the method of the present invention can be used in conjunction with those earlier selection methods should the need arise - if for example it is desirable to prepare cells that have been or are to be transformed with a number of NOIs.

Also, the selection method of the present invention can be used in conjunction with one or more other known selection methods, such as those that are described in WO 93/05163 (the contents of which are incorporated herein by reference) and/or WO 94/20627 (the contents of which are incorporated herein by reference) and/or PCT/GB98/00367 (the contents of which are incorporated herein by reference), should the need arise - if for example it is desirable to prepare cells that have been transformed with a number of NOIs.

In addition, the selection method of the present invention can be used in conjunction with one or more other selection methods according to the present invention should the need arise - if for example it is desirable to prepare cells that have been transformed with a number of NOIs.

A further beneficial use of a combination of selection methods according to the present invention results in a very efficient multiple screening technique. In this regard, and by way of example, the medium in the first screen utilising the selection method of the present invention would contain added low amounts of the component or the derivative thereof. With this first screen, selectable transformed cells are selected over at least the majority of the non-transformed cells. Then should - for example - any non-transformed cells be accidentally be carried over in that first screen then a second screen can be carried out. In the second screen the selected population of cells are subjected to a second selection method according to the present invention but wherein the component or the derivative thereof is present in the medium in a high concentration. In the second screen, predominantly the transformed cells would remain viable.

With this combined aspect of the present invention, the population of cells of the earlier aspects of the present invention can therefore be a pre-selected (e.g. pre-screened) population of cells, wherein the population of cells has been prior selected by one or more selection methods, such as those according to the present invention.

This combined aspect of the present invention can be alternatively expressed as: a selection method for selecting from a population of cells one or more selectable genetically transformed cells, wherein the population of cells comprises selectable genetically transformed cells and possible non-transformed cells; wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product; optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product; wherein a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product; wherein the component can be present in an amount in a medium

that is toxic to the non-transformed cells; the method comprising (a) the step of introducing the population of cells to a medium; wherein the component is or will be in such an amount that is utilisable by the transformed cells but in an amount that is toxic to the non-transformed cells; wherein each of the first expression product and the optional
5 second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof; and (b) a subsequent step of introducing at least portion of the transformed cells to a medium that
10 comprises a higher concentration of the component or the derivative thereof or precursor thereof.

The present invention also encompasses compositions and kits useful for this combined aspect of the present - such as the nucleotide sequence or expression product thereof
15 according to the present invention, a first medium containing no component or derivative thereof, and a second medium comprising a high concentration of the component or derivative thereof.

Furthermore, the selection method of the present invention can be used in conjunction
20 (such as sequentially) with further selection methods wherein those further selection methods are a combination of one or more other selection methods according to the present invention and one or more known selection methods - such as those that are dependent on antibiotic or herbicide resistance and/or those that are disclosed in WO 93/05163 and/or WO 94/20627.

25 In the selection methods of WO 93/05163 and/or WO 94/20627, the *manA* gene from *Escherichia coli*, which encodes mannose-6-phosphate isomerase (E.C. 5.3.2.8.), was employed as a selectable marker. This selection marker is suitable for *inter alia* the transformation of *Solanum tuberosum*, conferring positive selection in the presence of
30 mannose. In more detail, the coding region of *manA* was ligated into a CaMV 35S expression cassette, and introduced into a binary vector for plant transformation

mediated by *Agrobacterium tumefaciens*. To allow comparison of kanamycin selection with selection on mannose, the vector also contained a gene for kanamycin resistance, *nptII*. In order to identify transformants, the construction also contained the *B*-glucuronidase histochemical marker, *uidA*. Stable integration of the *manA* gene was shown by Southern blotting. Extracts from plants transformed with this construct, and selected on mannose, were shown to have specific activities for mannose-6-phosphate isomerase some five hundred fold those of control plants. Expression of *manA* in transformed cells relieved the metabolic paralysis, usually caused by mannose, while also allowing it to serve as a source of carbohydrate for transformants. These effects combined to impose a stringent selection pressure in favour of transformed cells, which allowed the recovery of transformants with a very low frequency of escapes. The percentage of shoots which were shown to be transgenic after selection on mannose was approximately twice that of shoots selected on kanamycin. The transformants selected on mannose have proven to be stable over three generations of plants propagated from tubers.

Hence, the population of cells of the earlier aspects of the present invention can therefore be a pre-selected (e.g. pre-screened) population of cells, wherein the population of cells has been prior selected by one or more selection methods according to the present invention and/or one or more other selection methods.

In addition, or in the alternative, the transformed cells selected by the selection method of the present invention can be subsequently subjected to one or more selection methods according to the present invention and/or one or more other selection methods.

The present invention also provides an expression system that enables transformed cells to be selected by the selection method of the present invention. The expression system can be expressing or can be capable of expressing at least the first nucleotide sequence of the present invention. The expression system may be one or more of a vector, construct, plasmid, cell or organism.

If a cell is also to be transformed with a NOI then the expression system will comprise that NOI - which NOI may be present on or in the same vector, construct, plasmid, cell or organism as the first nucleotide sequence. Alternatively the NOI may be present on or in a different vector, construct, plasmid, cell or organism as the first nucleotide sequence. Preferably, the NOI is present on or in the same vector, construct, plasmid, cell or organism as at least the first nucleotide sequence.

If a cell is to be transformed with one or more NOIs and one or more other genes for one or more other selection methods (such as another selection method according to the present invention and/or a known selection method) those other nucleotide sequences may be present on or in the same vector, construct, plasmid, cell or organism as the first nucleotide sequence. Alternatively one or more of those other nucleotide sequences may be present on or in a different vector, construct, plasmid, cell or organism as at least the first nucleotide sequence. Preferably, those other nucleotide sequences are present on or in the same vector, construct, plasmid, cell or organism as the first nucleotide sequence. This allows for workers to easily prepare and easily select for cells that have been transformed with a number of NOIs etc.

In accordance with the present invention there may be a plurality of NOI(s).

The, or each, NOI may be independently selected from DNA or RNA. Preferably, the or each NOI is DNA. More preferably, the or each NOI is recombinant DNA.

As indicated above, the term "recombinant DNA" means DNA prepared by at least one step that utilises at least one recombinant DNA technique.

The term NOI means any desired nucleotide sequence for incorporation into the cells in question to produce genetically transformed cells. Introduction of nucleotide sequences into, for example, plants is widely practised, and it is believed that there are no limitations upon the nucleotide sequences whose presence may be selected (eg. detected) by use of the selection method of the present invention.

By use of the method of the present invention the presence of the NOI in the genetically transformed cells may be determined without the above-mentioned disadvantages associated with the selection systems relying solely on antibiotic resistance and/or
5 herbicide resistance.

The NOI can be any nucleotide sequence of interest, such as any gene of interest. A NOI can be any nucleotide sequence that is either foreign or natural to the cell or organism (e.g. a particular plant) in question. Typical examples of a NOI include genes
10 encoding proteins and enzymes that modify metabolic and catabolic processes. The NOI may code for an agent for introducing or increasing resistance to pathogens. The NOI may even be an antisense construct for modifying the expression of natural transcripts present in the relevant tissues. The NOI may even code for a compound that is of benefit to animals or humans. Examples of NOIs include nucleotide sequences encoding
15 any one or more of pectinases, pectin depolymerases, polygalacturonases, pectate lyases, pectin lyases, rhamno-galacturonases, hemicellulases, endo- β -glucanases, arabinases, or acetyl esterases, or combinations thereof, as well as antisense sequences thereof. The NOI may encode a protein giving nutritional value to a food or crop. Typical examples include plant proteins that can inhibit the formation of anti-nutritive factors and plant
20 proteins that have a more desirable amino acid composition (e.g. a higher lysine content than a non-transgenic plant).

The NOI may even code for an enzyme that can be used in food processing such as chymosin, thaumatin and α -galactosidase. The NOI can be a gene encoding for any one
25 of a pest toxin, an antisense transcript such as that for patatin or α -amylase, ADP-glucose pyrophosphorylase (e.g. see EP-A-0455316), a protease antisense, a glucanase or genomic β 1,4-endoglucanase.

The NOI may even code for or comprise an intron of a particular gene. Here the intron
30 can be in sense or antisense orientation. In the latter instance, the particular gene could be DNA encoding β -1,4-endoglucanase. Antisense expression of genomic exon or intron

sequences as the NOI would mean that the natural β -1,4-endoglucanase expression would be reduced or eliminated but wherein the expression of a β -1,4-endoglucanase gene according to the present invention would not be affected.

5 The NOI may be the nucleotide sequence coding for the arabinofuranosidase enzyme which is the subject of PCT patent application PCT/EP96/01009. The NOI may be any of the nucleotide sequences coding for the ADP-glucose pyrophosphorylase enzymes which are the subject of PCT patent application PCT/EP94/01082. The NOI may be any of the nucleotide sequences coding for the α -glucan lyase enzyme which are described in
10 PCT patent application PCT/EP94/03397. The NOI may be any of the nucleotide sequences coding for the glucanase enzyme which are described in PCT patent application PCT/EP96/01008.

The NOI may also encode a permease or other transport factor which allows the
15 compound or precursor thereof or metabolised derivative thereof to cross the cell membrane and enter the transformed cells. Instead of facilitating uptake of a compound into a cell, the co-introduced nucleotide sequence may alternatively direct the component or precursor thereof or metabolised derivative thereof to a specific compartment - such as the plasma membrane or into the vacuole or the endoplasmic reticulum.

20 More than one NOI can be present.

The NOI can be co-introduced with the first nucleotide sequence according to the present invention.

25 The term "co-introduced" means that the two nucleotide sequences may be coupled to each other, or are otherwise introduced together, in such a manner that the presence of the co-introduced first nucleotide sequence in a cell indicates that the NOI has been introduced into the cell, i.e. if the first nucleotide sequence is shown to have been
30 introduced, the probability that the NOI has also been introduced is significantly

increased. The two nucleotide sequences may be part of the same genetic construct and may be introduced by the same vector.

The methods described herein may also be used when the co-introduced first nucleotide sequence and the NOI are introduced independently. This may be performed, for example, by using the same bacteria for incorporation of both genes and incorporating a relatively large number of copies of the NOI into the cells, whereby the probability is relatively high that cells which are shown to express the first nucleotide sequence will also contain and express the NOI.

In order for the introduced first nucleotide sequence and optional NOI to be expressed in the transformed cells, the genetic constructs containing the first nucleotide sequence and/or NOI will typically, but not necessarily, contain regulatory sequences enabling expression of the nucleotide sequences, e.g. known promoters and transcription terminators. Thus, the first nucleotide sequence will typically be associated with a promoter, which may be a constitutive or regulatable promoter, and the NOI will typically also be associated with a constitutive or regulatable promoter.

The term "promoter" is used in the normal sense of the art, e.g. an RNA polymerase binding site in the Jacob-Monod theory of gene expression.

The promoter could additionally include one or more features to ensure or to increase expression in a suitable host. For example, the features can be conserved regions such as a Pribnow Box or a TATA box. The promoters may even contain other sequences to affect (such as to maintain, enhance, decrease) the levels of expression of the nucleotide sequence of the present invention. For example, suitable other sequences include the *Shl*-intron or an ADH intron. Other sequences include inducible elements - such as temperature, chemical, light or stress inducible elements.

Also, suitable elements to enhance transcription or translation may be present. An example of the latter element is the TMV 5' signal sequence (see Sleat Gene 217 [1987] 217-225; and Dawson Plant Mol. Biol. 23 [1993] 97).

5 The first nucleotide sequence and/or the second nucleotide sequence and/or the third nucleotide sequence and/or the NOI may comprise one or more introns. In particular, if the first nucleotide sequence and/or the second nucleotide sequence and/or the third nucleotide sequence and/or the NOI encodes an expression product that can detrimentally affect a bacterium and all or a part (e.g. a plasmid thereof or therein) of that bacterium is
10 used either to propagate the NOI or as a means to transform the cells, then it may be desirable for that gene product to be inactive in the bacterium. One way of selectively inactivating the gene product in bacteria is to insert one or more introns into the nucleotide sequence of the first nucleotide sequence and/or the second nucleotide sequence and/or the third nucleotide sequence and/or the NOI, respectively. This intron
15 or those introns would not be removed after transcription in the bacterium but would be so removed in, for example, plants etc.

In a highly preferred embodiment, if the first nucleotide sequence and/or the second nucleotide sequence and/or the third nucleotide sequence and/or the NOI comprises at
20 least one intron, then that at least one intron is present in a highly conserved region of the first nucleotide sequence or the NOI.

Here, the term "intron" is used in its normal sense as meaning a nucleotide sequence lying within a coding sequence but being removable therefrom.

25

As mentioned above, the method of the present invention is particularly suitable for the selection of genetically transformed plant cells, thereby allowing identification and isolation of such cells without being essentially dependent on the use of selection genes coding for antibiotic or herbicide resistance.

30

The selection method of the present invention may be used for selecting cells *in vitro*. However, the selection method of the present invention may also be employed *in vivo* in the sense that it is possible to selectively grow transformed organisms - such as plants - from cells, tissues etc. that comprise the selection system of the present invention.

5 *In vivo* use of the selection method of the present invention is of particular importance in connection with genetic transformation performed on whole plants or on plant parts, in which the plants or plant parts comprise both transformed and non-transformed cells, since selection of the transformed cells can, in some instances, be achieved without
10 directly damaging the neighbouring non-transformed cells. For example, in some instances, the transformed cells have a selective advantage compared to the non-transformed cells - such as the ability to still form shoots - but the non-transformed cells suffer in the sense of being damaged or killed, as is the case with using antibiotics or herbicides.

15 In certain cases, such as when an improved selection frequency is desired, it may be advantageous for the cells to be transformed with a nucleotide sequence that is a selection gene different to the first nucleotide sequence. This additional, selection nucleotide sequence may be an additional gene coding for an enzyme (or other protein or
20 polypeptide) suitable for selection according to the present invention, or it may be a gene coding for an enzyme (or other protein or polypeptide) for a known selection method, eg coding for resistance to a antibiotic or herbicide or it may be a gene suitable for selection by the selection methods described in WO 93/05163 and/or WO 94/20627. Thus, genetically transformed cells may be selected using a combination of selection
25 techniques. For example, if the transformed cells also possessed genes coding for resistance to at least one antibiotic or herbicide, then the medium could additionally comprise at least one antibiotic or herbicide to which the transformed cells are resistant. In particular, we have found that the medium of the present invention does not impair the effectiveness of the known selection methods that rely on herbicide or antibiotic
30 resistance.

The selective advantage possessed by the transformed cells of the present invention may be any difference or advantage with regard to the non-transformed cells which allows the transformed cells to be readily identified and isolated from the non-transformed cells. This may, for example, be a difference or advantage allowing the transformed cells to be
5 identified by simple visual means, i.e. without the use of a separate assay to determine the presence of a gene that provides the selection means.

As mentioned above, one aspect of the present invention relates to genetically transformed cells which have been selected according to the above method, in particular
10 plant cells, as well as plants, progeny or seeds derived from or derivable from such genetically transformed plant cells. In particular, it is often an advantage that these cells are genetically transformed plant cells whose genome does not contain an introduced (i.e. non-native) nucleotide sequence coding for toxin-resistance, antibiotic-resistance or herbicide-resistance as a selection means. As explained above, there are concerns about
15 whether it is safe to incorporate genes coding for eg antibiotic resistance in eg food plants. Genetically transformed plant cells selected by the method of the present invention which do not contain selection genes for eg antibiotic resistance, as well as plants, progeny and seeds derived from such cells, are therefore clearly advantageous in this respect.

20

The transformed cells may be prepared by techniques known in the art. For example, if the transformed cells are transformed plant cells reference may be made to EP-B-0470145 and CA-A-2006454.

25 Even though the selection method according to the present invention is not disclosed in EP-B-0470145 and CA-A-2006454, those two documents do provide some useful background commentary on the types of techniques that may be employed to prepare the transformed plant cells and transgenic plants according to the present invention. Some of these background teachings are now included in the following commentary.

30

The basic principle in the construction of genetically modified plants is to insert genetic information in the plant genome so as to obtain a stable maintenance of the inserted genetic material.

5 Several techniques exist for inserting the genetic information, the two main principles being direct introduction of the genetic information and introduction of the genetic information by use of a vector system. A review of the general techniques may be found in articles by Potrykus (Annu Rev Plant Physiol Plant Mol Biol [1991] 42:205-225) and Christou (Agro-Food-Industry Hi-Tech March/April 1994 17-27).

10 Thus, in one aspect, the present invention relates to a vector system which carries a first nucleotide sequence or construct according to the present invention and which is capable of introducing the nucleotide sequence or construct into the genome of an organism, such as a plant.

15 The vector system may comprise one vector, but it can comprise at least two vectors. In the case of two vectors, the vector system is normally referred to as a binary vector system. Binary vector systems are described in further detail in Gynheung An et al. (1980), Binary Vectors, *Plant Molecular Biology Manual A3*, 1-19.

20 One extensively employed system for transformation of plant cells with a given promoter or nucleotide sequence or construct is based on the use of a Ti plasmid from *Agrobacterium tumefaciens* or a Ri plasmid from *Agrobacterium rhizogenes* (An et al. (1986), *Plant Physiol.* 81, 301-305 and Butcher D.N. et al. (1980), *Tissue Culture Methods for Plant Pathologists*, eds.: D.S. Ingrams and J.P. Helgeson, 203-208).

25 Several different Ti and Ri plasmids have been constructed which are suitable for the construction of the plant or plant cell constructs described above.

The first nucleotide sequence or construct of the present invention should preferably be inserted into the Ti-plasmid between the border sequences of the T-DNA or adjacent a T-DNA sequence so as to avoid disruption of the sequences immediately surrounding the T-DNA borders, as at least one of these regions appear to be essential for insertion of modified T-DNA into the plant genome.

As will be understood from the above explanation, if the organism is a plant, then the vector system of the present invention is preferably one which contains the sequences necessary to infect the plant (e.g. the *vir* region) and at least one border part of a T-DNA sequence, the border part being located on the same vector as the genetic construct. Preferably, the vector system is an *Agrobacterium tumefaciens* Ti-plasmid or an *Agrobacterium rhizogenes* Ri-plasmid or a derivative thereof, as these plasmids are well-known and widely employed in the construction of transgenic plants, many vector systems exist which are based on these plasmids or derivatives thereof.

In the construction of a transgenic plant the promoter or nucleotide sequence or construct of the present invention may be first constructed in a microorganism in which the vector can replicate and which is easy to manipulate before insertion into the plant. An example of a useful microorganism is *E. coli.*, but other microorganisms having the above properties may be used. When a vector of a vector system as defined above has been constructed in *E. coli.* it is transferred, if necessary, into a suitable *Agrobacterium* strain, e.g. *Agrobacterium tumefaciens*. The Ti-plasmid harbouring the first nucleotide sequence or construct of the invention is thus preferably transferred into a suitable *Agrobacterium* strain, e.g. *A. tumefaciens*, so as to obtain an *Agrobacterium* cell harbouring the promoter or nucleotide sequence or construct of the invention, which DNA is subsequently transferred into the plant cell to be modified.

Naturally, the present invention is not limited to just the use of *Agrobacterium* systems to transform plants. In this regard, other suitable techniques may be used - such as electroporation and/or particle bombardment (biolistics).

As reported in CA-A-2006454, a large number of cloning vectors are available which contain a replication system in *E. coli* and a selection means which allows a selection of the transformed cells. The vectors contain for example pBR322, the pUC series, the M13 mp series, pACYC 184 etc. In this way, the promoter or nucleotide or construct of the present invention can be introduced into a suitable restriction position in the vector. The contained plasmid is used for the transformation in *E. coli*. The *E. coli* cells are cultivated in a suitable nutrient medium and then harvested and lysed. The plasmid is then recovered and then analysed - such as by any one or more of the following techniques: sequence analysis, restriction analysis, electrophoresis and further biochemical-molecular biological methods. After each manipulation, the used DNA sequence can be restricted or selectively amplified by PCR techniques and connected with the next DNA sequence. Each sequence can be cloned in the same or different plasmid.

After each introduction method of the first nucleotide sequence or construct according to the present invention in the plants the presence and/or insertion of further DNA sequences may be necessary. If, for example, for the transformation the Ti- or Ri-plasmid of the plant cells is used, at least the right boundary and often however the right and the left boundary of the Ti- and Ri-plasmid T-DNA, as flanking areas of the introduced genes, can be connected. The use of T-DNA for the transformation of plant cells has been intensively studied and is described in EP-A-120516; Hoekema, in: The Binary Plant Vector System Offset-drukkerij Kanters B.B., Alblaserdam, 1985, Chapter V; Fraley, *et al.*, Crit. Rev. Plant Sci., 4:1-46; and An et al., EMBO J. (1985) 4:277-284.

Direct infection of plant tissues by *Agrobacterium* is a simple technique which has been widely employed and which is described in Butcher D.N. et al. (1980), *Tissue Culture Methods for Plant Pathologists*, eds.: D.S. Ingrams and J.P. Helgeson, 203-208. For further teachings on this topic see Potrykus (Annu Rev Plant Physiol Plant Mol Biol [1991] 42:205-225) and Christou (Agro-Food-Industry Hi-Tech March/April 1994 17-

27). With this technique, infection of a plant may be done on a certain part or tissue of the plant, i.e. on a part of a leaf, a root, a stem or another part of the plant.

Typically, with direct infection of plant tissues by *Agrobacterium* carrying the first nucleotide sequence or the construct, a plant to be infected is wounded, e.g. by cutting the plant with a razor or puncturing the plant with a needle or rubbing the plant with an abrasive. The wound is then inoculated with the *Agrobacterium*. The inoculated plant or plant part is then grown on a suitable culture medium.

When plant cells are constructed, these cells are grown and, optionally, maintained in a medium according to the present invention following well-known tissue culturing methods - such as by culturing the cells in a suitable culture medium supplied with the necessary growth factors such as amino acids, plant hormones, vitamins, etc, but wherein the culture medium comprises a component according to the present invention.

Regeneration of the transformed cells into genetically modified plants may be accomplished using known methods for the regeneration of plants from cell or tissue cultures, for example by selecting the transformed shoots and by subculturing the shoots on a medium containing the appropriate nutrients, plant hormones, etc.

Further teachings on plant transformation may be found in EP-A-0449375.

Reference may even be made to Sprngstad *et al* (1995 Plant Cell Tissue Organ Culture 40 pp 1-15) as these authors present a general overview on transgenic plant construction.

In a highly preferred embodiment, the present invention is based on our finding that it is possible to use constructs comprising an expressable nucleotide sequence gene coding for any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) to prepare transformed cells wherein the transformed cells can be selected from non-transformed cells.

In addition, the present invention also covers transgenic plants comprising the transformed cells or constructs of the present invention.

Thus, in a highly preferred embodiment the present invention covers transgenic plants comprising transformed cells or constructs that comprise an expressable nucleotide sequence gene coding for any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2).

In order to explain in more detail these highly preferred aspects of the present invention, reference shall be made to at least Figures 1 - 2, in which

Figure 1 is a schematic diagram of a metabolic pathway;

Figure 2 is a schematic diagram of a metabolic pathway.

Galactose metabolism in plants have been studied in *Petunia* which is one of the very few plant species being able to use galactose as carbon source to sustain growth (Dressler *et al.* 1982). The first step of galactose metabolism in *Petunia* as well as in yeast and other microorganisms is the phosphorylation of galactose to galactose-1-phosphate by galactokinase (EC 2.7.1.6) (see Figs. 1 and 2 below). The next step is the conversion to UDP-galactose which in *Petunia* can be performed by either a UTP-dependent pyrophosphorylase (EC 2.7.7.10) or a UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12) while in microorganisms only or predominantly the latter reaction occurs. The UDP-galactose is finally converted to UDP-glucose by UDP-galactose epimerase (EC 5.1.3.2). Thus, the conversion of galactose to a glucose-containing metabolite is irreversible.

The reason for the toxicity of galactose on plant cells is believed to be due to the absence of one or more of the enzymatic activities required for the conversion of galactose to UDP-glucose. Thus, in order to use galactose as selective agent, the transformed plant

cells must be supplemented with the lacking gene(s) so that enzymatic activities for the entire pathway from galactose to UDP-glucose are present.

However, it is likely that many plant species do not require the insertion of all 3 genes in the Leloir pathway as the absence of just one gene in this pathway will render the plant tissue sensitive to galactose.

The first enzyme in galactose metabolism is galactokinase which has been reported to be present in some plants such as *Vicia faba* (Dey 1983), *Phaseolus aureus* (Chan and Hassid 1975; Neufeld *et al.* 1960), barley and corn (Roberts *et al.* 1971). Roberts *et al.* (1971) studied the accumulation of galactose containing metabolites in corn roots fed with toxic levels of galactose and they found that free galactose and galactose-1-phosphate each accounted for close to 50% of the indentifiable metabolites. These data were in accordance with the hypothesis that the causal agent of galactose toxicity may be galactose-1-phosphate. As galactose is toxic to most plant species, this in turn implies that many plant species have galactokinase activity.

The next enzyme in the galactose metabolism is UTP-hexose-1-phosphate uridylyltransferase or UDP-glucose uridylyltransferase. Both of these enzymes - which are rare - may be found in *Petunia*.

In contrast to the two preceeding enzymes, the last enzyme, UDP-galactose epimerase, has been reported in a range of plant species such as fenugreek (Clermont and Percheron 1979), wheat (Fan and Feingold 1969), sugarcane (Maretzki and Thom 1978) and various trees (Dalessandro and Northcote 1977) in addition to the above mentioned species. The reason for the widespread occurrence of this enzyme is presumably related to the fact that its main function is the production of UDP-galactose from UDP-glucose for the synthesis of cell wall components.

Thus, in order to enable transformed plant tissue to metabolise galactose, it is likely that it is sufficient for some plant species to introduce just one gene eg. a UTP-hexose-1-phosphate transferase gene or a UDP-glucose uridylyltransferase gene which would then function as the selectable gene. However, for other plant species it may be required to
5 introduce two or even three genes involved in the metabolism of galactose. From a technical point of view, it presents no problems to use more than one gene as selectable gene.

Although a galactose selection system also employs a carbohydrate as selective agent, it
10 does not resemble 'positive selection' due to the high toxic effect of galactose.

The selection system described here is based on enabling transformed plant cells to metabolise D-galactose. This D-galactose can be supplied to the plant cells or tissues during selection as free D-galactose or as a D-galactose containing compound. Examples
15 of such D-galactose containing compounds are α -lactose (β -D-galactopyranosyl [1 \rightarrow 4]- α -D-glucose, milk sugar), β -lactose, melibiose (6-O- α -D-galacto-pyranosyl-D-glucose), raffinose, stachyose, verbascose and galactinol which all liberate free D-galactose upon hydrolysis by either α -galactosidases or β -galactosidases which are present in a wide range a plant species and tissues.

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Other examples of potentially useful compounds for galactose selection are chemically derivatized D-galactose such as D-galactose pentaacetate and methyl galactoside, from which free D-galactose can be produced by the action of appropriate enzymes present in plant cells.

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It can also envisaged that the galactose selection system can be modified so that galactose-1-phosphate can be used as selective agent, in which case the selectable gene is most likely to be an UTP-hexose-1-phosphate uridylyltransferase gene or an UDP-glucose uridylyltransferase gene.

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5 Yet another modification of the galactose selection system relates to the addition of other carbohydrates or other substances to the selection medium which affect the toxicity of galactose. Furthermore, it will also be obvious for those skilled in the art that changes in the employed galactose concentration in the course of the selection can affect selection efficiency.

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Thus, the present invention provides a new selection method based on the novel use of galactose and/or a derivative thereof and/or a precursor thereof.

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As galactose and the genes encoding the enzymes responsible for the metabolism of galactose are well-characterized, we believe that galactose would be suitable for use as a selective agent for the selection of transgenic cells, tissues, organs, organisms (such as plants), provided that the transgenic cells, tissues, organs, organisms (such as plants) acquire the appropriate genes for metabolising, detoxifying and/or tolerating galactose.

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The present invention will now be described only by way of examples in which reference is made to the following Figures:

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Figure 3 is presents sequence information; and

Figure 4 is a schematic diagram of a plasmid.

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In more detail, Figure 3 is the complete nucleotide sequence of the *E. coli galT* gene with the deduced amino acid sequence shown below the nucleotide sequence.

In more detail, Figure 4 is a plasmid map of the plant transformation vector pVIC-GalT.

TRANSFORMATION STUDIES

The following examples demonstrate that the genes coding for galactokinase (EC 2.7.1.6) (*galK*), UTP-dependent pyrophosphorylase (EC 2.7.7.10) (*galP*), UDP-glucose-dependent uridyl transferase (EC 2.7.7.12) (*galT*), UDP-galactose epimerase (EC 5.1.3.2) (*galE*) gene can be used as a means to provide selection of transformed cells, such as transgenic potato or maize shoots, on or in media containing galactose or a derivative thereof as a selective agent. For convenience, the term *galX* has been used to denote any one or more of *galK*, *galP*, *galT*, *galE*.

EXAMPLE 1 - TRANSGENIC POTATO PLANTS

General teachings on potato transformation may be found in our copending patent applications PCT/EP96/03053, PCT/EP96/03052 and PCT/EP94/01082 (the contents of each of which are incorporated herein by reference).

For the present studies, the following protocol was adopted.

Plasmid construction

The disarmed *Agrobacterium tumefaciens* strain LBA 4404, containing the helper *vir* plasmid pRAL4404 (Hoekema *et al*, 1983 Nature 303 pp 179-180), was cultured on YMB agar ($K_2HPO_4 \cdot 3H_2O$ 660 mg l⁻¹, $MgSO_4$ 200 mg l⁻¹, NaCl 100 mg l⁻¹, mannitol 10 g l⁻¹, yeast extract 400 mg l⁻¹, 0.8% w/v agar, pH 7.0) containing 100 mg l⁻¹ rifampicin and 500 mg l⁻¹ streptomycin sulphate. Transformation with a plasmid containing *galX* under the control of a plant expressible promoter (such as the E35S promoter) was accomplished using the freeze-thaw method of Holters *et al* (1978 Mol Gen Genet 163 181-187) and transformants were selected on YMB agar containing 100 mg l⁻¹ rifampicin and 500 mg l⁻¹ streptomycin, and 50 mg l⁻¹ gentamycin sulphate. The resultant plasmid is called pVICTOR IV 35S *galX*. Transformation with a control construct lacking the *galX* gene was performed in the same manner. In addition, the T-DNA will include a

screenable marker gene encoding a screenable marker enzyme, such as the β -*glucuronidase* gene from *E. coli*, driven by a plant expressable promoter. This facilitates the optimization of the methods as transgenic cells/shoots/plants are more easily found using this marker (Jefferson *et al* 1987, EMBO J, 6:3901-3907).

5

Transformation of plants

Shoot cultures of *Solanum tuberosum* cv Saturna were maintained on LS agar containing Murashige Skoog basal salts (Sigma M6899) (Murashige and Skoog (1965) Physiol. Plant. 15: 473-497) with 2 μ M silver thiosulphate, and nutrients and vitamins as described by Linsmaier and Skoog (1965 Physiol. Plant. 18 100-127). Cultures were maintained at 25°C with a 16 h daily photoperiod. After approximately 40 days, subculturing was performed and the shoots cut into segments of approximately 8 mm length.

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Shoot cultures of approximately 40 days maturity (5-6 cm height) were cut into 8 mm internodal segments and/or leaves were cut off and wounded by making 2-4 small cuts over the midrib of the leaf. These were then placed into liquid LS-medium containing *Agrobacterium tumefaciens* transformed with pVICTOR IV 35S *galX* ($A_{660} = 0.5$, pathlength 1 cm). Following incubation at room temperature for 30 minutes, the segments were dried by blotting on to sterile filter paper and transferred to LS agar (0.8% w/v containing 2 mg l⁻¹ 2,4-dichlorophenoxyacetic acid and 500 μ g l⁻¹ trans-zeatin. The explants were covered with filter paper, moistened with LS medium, and covered with a cloth for three days at 25°C. Following this treatment, the segments can be washed with liquid LS medium containing 800 mg l⁻¹ carbenicillin, and then transferred on to LS agar (0.8% w/v) containing 1 mg l⁻¹ trans-zeatin, 100 mg l⁻¹ gibberellic acid (GA3), with sucrose (eg 10-20 g l⁻¹). This agar contains galactose (eg such as in an amount of from about 0.5 - 5.0 g l⁻¹).

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The segments were sub-cultured to fresh substrate each 3-4 weeks. In 3 to 4 weeks, shoots develop from the segments and the formation of new shoots continued for 3-4 months.

- 5 The regenerated shoots are maintained on substrate composed of LS-substrate, 0.002 mM silver thiosulphate and agar (8.0 g/l). Carbenicillin (800 mg/l) can be added if desired.

- 10 The transgenic plants may be verified by performing a β -glucuronidase assay on the leaf tips of the surviving shoots according to Hodal *et al.* (Plant. Sci. (1992), 87: 115-122).

Alternatively, the transgenic genotype of the regenerated shoot may be verified by performing NPTII assays (Radke, S. E. *et al.*, Theor. Appl. Genet. (1988), 75: 685-694) or by performing PCR analysis according to Wang *et al.* (1993, NAR 21: 4153-4154).

- 15 The shoots (height approximately 2-3 cms) were transplanted from rooting substrate to soil and placed in a growth chamber (21°C, 16 hour light 200-400 μ E/m²/sec).

- 20 When the plants were well established they were transferred to the greenhouse, where they were grown until tubers had developed and the upper part of the plants were senescing.

Harvesting

- 25 The potatoes were harvested after about 3-6 months and then analysed.-

- 30 The transformed shoots can be distinguished from the non-transformed shoots by adding galactose to their substrate of leaf tips cultured in in-vitro. After harvest of the shoots, the transformed shoots can be selected by adding amounts of galactose to the shoot medium. The transformed shoots will be resistant to galactose and will survive as opposed the non-transformed shoots which will be inhibited in growth.

Analysis of Transformants

In order to confirm the integration of *galX*, genomic DNA may be isolated by the method of Dellaporta *et al* (1983 Plant Mol Biol Rep 1 19-21) and samples of this DNA, digested with *EcoRI*, subjected to electrophoresis in an 0.8% w/v agarose gel and transferred to Hybond N+ membranes (Amersham) by Southern blotting (Southern, 1975 J Mol Biol 98 503-517). Probes for the coding region of *galX* may be used as templates for random primed synthesis of ³²P-labelled probe after the method of Feinberg and Vogelstein (1983 Anal Bioch 137 266-267) and hybridised to the Southern blots at high stringency (65°C, 0.1 x SSC).

Selection of transgenic shoots was accomplished using a selection medium according to the present invention.

Transgenic shoots are obtained on selection media according to the present invention indicating that the selection medium can be varied significantly and remain useful for the selection of transgenic shoots.

EXAMPLE 2 - TRANSGENIC MAIZE PLANTS

Introduction

Since the first publication of production of transgenic plants in 1983 (Leemans, 1993 Biotechnology 11 s22), there have been numerous publications of production of transgenic plants including especially dicotyledon crop plants.

Until very recently there were very few reports on successful production of transgenic monocotyledonary crop plants. This relatively slow development within monocots were due to two causes. Firstly, until the early 1980s, efficient regeneration of plants from cultured cells and tissues of monocots had proven very difficult. This problem was

ultimately solved by the culture of explants from immature and embryogenic tissue, which retain their morphogenic potential on nutrient media containing plant growth regulators. Secondly, the monocots are not a natural host for *Agrobacterium tumefaciens*, meaning that the successful developed techniques within the dicots using their natural vector *Agrobacterium tumefaciens* was unsuccessful for many years in the monocots.

Nevertheless, it is now possible to successfully transformation and produce fertile transgenic plants of maize using methods such as: (1) Silicon Carbide Whiskers; (2) Particle Bombardment; (3) DNA Uptake by PEG treated protoplast; or (4) DNA Uptake in Electroporation of Tissue. Each of these methods - which are reviewed by Thompson (1995 Euphytica 85 pp 75-80) - may be used to prepare *inter alia* transgenic maize according to the present invention.

In particular, the particle Gun method has been successfully used for the transformation of monocots. However, EP-A-0604662 reports on a different method of transforming monocotyledons. The method comprises transforming cultured tissues of a monocotyledon under or after dedifferentiation with *Agrobacterium* containing a super binary vector as a selection means a hygromycin-resistant gene was used. Production of transgenic calli and plant was demonstrated using the hygromycin selection. This method may be used to prepare *inter alia* transgenic maize according to the present invention.

Subsequent to the method of EP-A-0604662, EP-A-0672752 reports on non-dedifferentiated immature embryos. In this regard, both hygromycin-resistance and PPT-resistance genes were used as the selection means, with PPT giving rise to 10% or more independent transformed plants. This method may be used to prepare *inter alia* transgenic maize according to the present invention.

To date, it would appear that transgenic maize plants can be successfully produced from easily-culturable varieties - such as the inbred line A188. In this regard, see the

teachings of Ishida *et al* (1996 Nature Biotechnology 14 pp 745-750). The method disclosed by these workers may be used to prepare *inter alia* transgenic maize according to the present invention.

- 5 Vasil (1996 Nature Biotechnology 14 pp 702-703) presents a further review article on transformation of maize.

Even though it is possible to prepare transformed maize by use of, for example, particle Gun mediated transformation, for the present studies the following protocol is adopted.

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Plasmid construction

The same protocol as outlined above is adopted. In this respect, the protocol also uses pVICTOR IV 35S *galX*. Likewise, transformation with a control construct lacking the *galX* gene was performed in the same manner.

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Isolation and cocultivation of explants

Immature embryos of, for example, maize line A188 of the size between 1.5 to 2.5 mm were isolated and cocultivated with *Agrobacterium tumefaciens* strain LBA 4404 in N6-AS for 2-3 days at 25°C under illumination. Thereafter, the embryos were washed with sterilized water containing 250 mg/l of cefotaxime and transferred to an LS medium and 250 mg/l cefotaxime and galactose in concentrations of up to 100 mg/l (the medium is hereafter called LSS2).

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Conditions for the selection of transgenic plants

The explants were cultured for three weeks on LSS2 medium and then transferred to an LS medium containing galactose and optionally cefotaxime. After three weeks on this medium, green shoots were isolated and tested for β -glucuronidase (Jefferson *et al* 1987, EMBO J 6:3901-3907) activity.

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Rooting of β -glucuronidase positive shoots

5 β -glucuronidase positive shoots were transferred to an MS medium containing 2 mg/l for rooting. After four weeks on this medium, plantlets are transferred to pots with sterile soil for acclimatisation.

Selection of transgenic shoots was accomplished using a selection medium according to the present invention.

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After four weeks the shoots were harvested and all explants were transferred to fresh selection medium (same composition) and after another four weeks of selection the last shoots were harvested.

15 After harvest, the shoots were analysed for GUS activity using the histochemical assay.

Transgenic shoots are obtained on selection media according to the present invention indicating that the selection medium can be varied significantly and remain useful for the selection of transgenic shoots.

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EXAMPLE 3 - TOXICITY OF GALACTOSE TO VARIOUS PLANT SPECIES

In order to be useful for selection, the selective component (otherwise called selective agent) should not be able to sustain significant growth and preferably have adverse/toxic effects to the non-transformed cells/tissues. To assess the applicability of galactose selection following transformation, a number of plant species representing a range of different botanical families were tested for sensitivity to galactose, using explants which may be suitable for transformation experiments.

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Galactose was unable to sustain growth of any of the tested plant species (wheat, sunflower, oil seed rape, potato, sugar beet, pea and Petunia) with varying levels of toxicity. The only fairly tolerant species was Petunia, in accordance with expectations as Petunia has been reported to have all necessary enzymatic activities for metabolising galactose (Dressler *et al.* 1982. Z. Pflanzenphysiol. 107: 409-418).

3.1. Wheat (*Triticum aestivum*)

As an example of a monocotyledonous species, wheat was studied. Immature wheat embryos (2-3 weeks old, a total of 161) were carefully removed from the seed and were cultivated on Gamborg B5 medium (Gamborg *et al.* 1968. Exp. Cell Res. 50: 151-158) supplemented with 9.0 g/l Bacto agar, 20 g/l sucrose and galactose according to the experiment. The embryos were cultivated in the dark for about 10 days where germinated embryos were transferred to fresh medium of the same composition and cultivated in a 16 h/8 h day/night regime for 4-5 weeks. The effect of galactose was evaluated by the number of germinated embryos and by measuring the length of the leaf on these embryos, relative to the control medium (no galactose).

Galactose concentration (g/l)	0	2.5	5.0	7.5	10	15
Germination of embryos	100	43	29	0	14	0
Average shoot length	100	89	67	0	33	0

The data in the above table clearly demonstrate that galactose has a strong inhibiting effect on the immature embryos as both germination and growth of the shoots were significantly reduced. The galactose selection method of the present invention can be used for the production of transgenic wheat.

Roberts *et al.* (1981. Plant Physiol. 48: 36-42) found that galactose was toxic to tissues of maize (*Zea mays*) and of barley (*Hordeum vulgare*). This indicates that the galactose selection method of the present invention may also be useful for the selection of transgenic shoots of these species.

3.2. Sunflower (*Helianthus annuus*)

The preparation of apical explants useful for the transformation of sunflower were modified according to Knittel *et al.* (1994. Plant Cell Rep. 14: 81-86). The seed coats of 2 d old seedlings were gently removed and the cotyledons, the emerging root and the top of the apex were removed. The remaining part of the apex was cut off and divided into two halves and cultivated according to Knittel *et al.*, with 20 g/l sucrose in the medium and galactose according to the experiment. After 4-5 weeks of culture at a 16 h/8 h day/night regime, growth was monitored by determining the fresh weight of the explants (above controls grown on a medium devoid of any carbohydrate source).

Galactose concentration (g/l)	0	5.0	7.5	10	15
Fresh weight (mg/explant)	144	28	0	0	0

The data show that galactose is toxic to sunflower explants. The data indicate that the galactose selection method of the present invention may be employed successfully for the selection of transgenic sunflower shoots.

3.3. Oil seed rape (*Brassica napus*)

Hypocotyl explants useful for the transformation of oil seed rape were prepared using a protocol modified after Radke *et al.* (1988. Theor. Appl. Genet. 78: 161-168). Hypocotyls of 5 d old seedlings were cut into 1 cm segments and transferred to modified

MS-medium (Murashige and Skoog 1962. *Physiol. Plant.* 15: 473-497) containing sucrose and galactose according to the experiment. After 3-4 weeks, shoots were harvested and the explants were transferred to fresh medium of the same composition. This was repeated for 3 more harvests and the total number of regenerated shoots formed on each medium was calculated (see table below).

Sucrose concentration (g/l)	0	5.0	10	20
0 g/l galactose	0	-	-	60
2.5 g/l galactose	0	1	7	26
5.0 g/l galactose	0	0	3	22

The data in the above Table clearly show that galactose is toxic to oil seed rape. Hence, the data indicate that the galactose selection method of the present invention may be used for the selection of transgenic shoots of oil seed rape and presumably related species. It is expected that higher concentrations of galactose are even more toxic.

The data also demonstrate that the toxic effect of galactose strongly interacts with sucrose as increasing concentrations of sucrose progressively reduces the toxic effect of galactose. Hence, for some applications, preferably galactose is present in low amounts of sucrose.

3.4. Potato (*Solanum tuberosum*)

Young leaves useful for the transformation of potato were prepared from leaves of 28 d old in vitro shoot cultures. The leaves were cut off avoiding the meristem at the leaf corner and were cut twice across the midvein. The explants were then put on modified MS-medium (Murashige and Skoog 1962. *Physiol. Plant.* 15: 473-497) containing various concentrations of sucrose and galactose. After 3 weeks, shoots were harvested and the explants were transferred to fresh medium of the same composition. This was repeated for 2 more harvests and the total number of explants with regenerated shoots formed on each medium was calculated (see table below).

Sucrose concentration (g/l)	0	5.0	10	20	30
0 g/l galactose	0	-	-	-	14
2.5 g/l galactose	1	0	0	7	-
5.0 g/l galactose	0	0	0	0	-

The data in the above Table clearly show that galactose is toxic to potato. The data indicate that the galactose selection method of the present invention may be used for the selection of transgenic shoots of potato and presumably related species. Like oil seed rape, these data also demonstrate that the toxic effect of galactose strongly interacts with sucrose as increasing concentrations of sucrose progressively reduces the toxic effect of galactose. We believe that such interaction is likely to take place also for many other plant species. We also believe that such interaction will also be observed with other carbohydrates than sucrose, such as glucose, fructose and maltose etc. Hence, for some applications, preferably galactose is present in low amounts of carbohydrate (i.e. in amounts that do not affect substantially the effect of the galactose).

3.5. Sugar beet (*Beta vulgaris*)

Cotyledonary explants useful for the transformation of sugar beet were prepared by
5 excising the cotyledons from 3-4 weeks old seedlings (Joersbo *et al.* 1998. Mol.
Breeding 4: 111-117). The leaves were transferred to modified MS-medium (Murashige
and Skoog 1962. Physiol. Plant. 15: 473-497) containing 20 g/l sucrose and various
concentrations galactose.

10 After 4-5 weeks, the number of shoots and the fresh weight of the explants were
determined and a significant reduction of both growth parameters was observed. The
data indicate that the galactose selection method of the present invention may be used for
the selection of transgenic shoots of sugar beet and presumably related species.

3.6. Pea (*Pisum sativum*)

Hypocotyl explants useful for the transformation of pea were prepared by excising the
hypocotyl from 7 d old seedlings and transferred to a medium, according to Nielsen *et*
al. (1991. Physiol. Plant 82: 99-102), containing various concentrations of sucrose and
20 galactose. After 5 weeks, the growth of the explants were evaluated. The results
showed that galactose was unable to sustain any growth of the explants, as opposed to
the control medium containing 20 g/l sucrose (no galactose) where the explants produced
a many-fold increased fresh weight. The data indicate that the galactose selection method
of the present invention may be used for the selection of transgenic shoots of pea and
25 presumably related species.

EXAMPLE 4 GALACTOSE SELECTION EMPLOYED FOR THE SELECTION OF TRANSGENIC SHOOTS

Introduction

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In one preferred embodiment, it is desirable to use up to 3 genes encoding enzymes useful for the conversion of galactose to UDP-glucose. However, in some instances we have found that only one gene may be used. An example of such a gene is that coding for a UTP-dependent pyrophosphorylase (EC 2.7.7.10) or a UDP-glucose-dependent uridyl transferase (EC 2.7.7.12). In some instances, these individual genes may be sufficient for the selection of transgenic shoots of many plant species because both galactokinase (EC 2.7.1.6) and UDP-galactose epimerase (EC 5.1.3.2) occur naturally in many plant species.

15 In this example, a further example (i.e. in addition to Example 1) the successful selection of transgenic potato shoots is described. These particular experiments used a transformation vector including an *E. coli* gene (*galT*) encoding UDP-glucose-dependent uridyl transferase (EC 2.7.7.12).

20 The fact that it was possible select transgenic potato shoots on galactose containing media is in accordance with the toxicological test where potato was found to be sensitive to galactose. Consequently, it is expected that transgenic shoots of other plant species that are sensitive to galactose also can be selected using galactose selection.

25 4.1. Transformation vector

An *E. coli galT* gene encoding galactose-1-phosphate uridylyltransferase has been cloned, sequenced and introduced into a plant transformation vector. The cloned gene consists of 1060 base pairs and codes for a 348 amino acid protein. Transformation of a *galT* deficient *E. coli* strain resulted in colonies able to grow on galactose as sole carbon source, indicating that the cloned gene expresses a functional enzyme capable of

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complementing the *galT* deficiency. The gene was furnished with an enhanced 35S promoter and a 35S terminator and inserted into a plant transformation vector based on our pVIC vector, into which a GUS gene was also inserted.

5 Cloning and characterization of the *E. coli galT* gene.

The *galT* gene of *E. coli* was isolated and cloned by PCR from genomic DNA isolated from an *E. coli* strain that can metabolise galactose. The *galT* gene was found in the GenBank Sequence Database, and the nucleotide sequence was used to design PCR
10 primers for the amplification of the coding region of the gene. To facilitate cloning into our expression vectors the upstream primer was designed such that a *Pst*I site was added to the 5' end of the primer, and the downstream primer was similarly designed with a *Sa*I site. A 1050 base pair DNA fragment obtained after PCR was cloned into the TA cloning vector pCR 2.1-TOPO. Five of the clones were completely sequenced and one
15 clone was selected for further use. The nucleotide sequence of the selected clone (Fig. 3) was identical to that of Cornwell *et al.* (1987. Nucl. Acids Res. 15: 8116). Translation of the cloned *galT* gene revealed an open reading frame of 1047 base pairs encoding a 348 amino acid protein (Fig. 3) in agreement with the published sequence (Cornwell *et al.* 1987. Nucl. Acids Res. 15: 8116).

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Construction of a plant transformation vector with the *galT* gene

A plant transformation vector called pVIC for *Agrobacterium* mediated transformation was used. pVIC contains the following features: Right- and left border sequences (RB
25 and LB) are from the *A. tumefaciens* plasmid pTiT37 (Yadav *et al.* 1982. Proc. Natl. Acad. Sci. 79: 6322-6326). The replication origin (pUC Ori) for replication and maintenance in *E. coli* is from the plasmid pUC19 (Yanish-Perron *et al.* 1985. Gene 33: 103-119). The replication origin (pVS1 Ori) for replication and maintenance in *A. tumefaciens* is from the *Pseudomonas* plasmid pVS1 (Itoh *et al.* 1984. Plasmid 11:
30 206-220). The bacterial spectinomycin/streptomycin resistance gene (Spec/Strep) is from the transposon Tn7 (Fling *et al.* 1985. Nucl. Acids Res. 19: 7095-7106) and is

fused to the hybrid trp-lac promoter (Amman *et al.* 1983. Gene 25: 167-178) for efficient bacterial expression. A β -glucuronidase gene (Jefferson *et al.* 1986. Proc. Natl. Acad. Sci. 83: 8447-8451) containing an intron of the ST-LS1 gene from potato, preventing expression in bacteria (Vancanneyt *et al.* 1990. Mol. Gen. Genet. 220: 245-250) was furnished with a 35S promoter and a 35S terminator and was introduced into pVIC. This construct was used as a vector for the *galT* gene. The cloned *galT* gene was cloned between an enhanced 35S promoter (E35S PR) and the 35S terminator (35S t) (Key *et al.* 1987. Science 236: 1299-1302 and Odell *et al.* 1985. Nature 313: 810-812) and this expression cassette was ligated into above mentioned vector. This resulted in the final plant transformation vector pVIC-galT, as shown schematically in Fig. 4.

4.2. Transformation of potato explants

Young leaves useful for the transformation of potato were prepared from leaves of 28 d old in vitro shoot cultures. The leaves were cut off avoiding the meristem at the leaf corner and were cut twice across the midvein. The explants were inoculated for 30 min. with a suspension of *Agrobacterium tumefaciens* strain EHA101 ($OD_{660}=0.5$) harbouring the transformation vector described above. After inoculation, the explants were dried carefully on filterpaper and left for co-cultivation for 3 d in dim light.

4.3. Selection of transgenic potato shoots on galactose containing media

After co-culture, the explants were transferred to selection medium consisting of MS-medium (Murashige and Skoog 1962. Physiol. Plant. 15: 473-497) supplemented with 500 mg/l carbenicillin and various concentrations of sucrose and galactose. After 3 and 6 weeks, shoots were harvested and transgenicity was evaluated by subjecting the harvested shoots to the assay for β -glucuronidase activity (Jefferson *et al.* 1987. Plant Mol. Biol. Rep. 5: 387-405), a marker enzyme encoded by a gene placed adjacent the selective gene on the T-DNA of the transformation vector (Fig. 4). The number of transgenic shoots isolated on the various selection media is indicated in the table below.

Galactose concentration (g/l)	Sucrose concentration (g/l)	No. of transgenic shoots	Transformation frequency (%)
0	20	0	0
1.25	10	3	6.0
1.25	15	7	14.0
1.25	20	3	6.0
2.5	10	1	2.0
2.5	15	0	0
2.5	20	4	8.0

The data show that it is possible to select transgenic shoots of a plant, transformed with a gene encoding an enzyme that enables or enhances the ability of the transformed shoots to reduce the toxicity of galactose.

In an other experiment, a few transgenic potato shoots (4%) were harvested on a medium devoid of galactose. Without wishing to be bound by theory we believe that this selective effect may be related to a stimulating effect of the inserted selective gene *per se* and/or the expression product thereof *per se* on the regeneration potential of the transgenic cells.

In combination with the toxicological studies, the fact that transgenic potato shoots can be selected on galactose-containing media suggests that transgenic shoots of other plant species that are unable to grow on or inhibited by galactose also may be selected.

- 5 Without wishing to be bound by theory, we also believe that the degree of selectivity may differ due to the choice of the selection media - which may contain galactose and/or metabolic precursors and/or metabolic derivatives and/or derivatives of galactose, alone or in combination with other carbohydrate/energy source(s) such as sucrose, glucose, fructose, maltose which may or may not affect the effect of galactose. By way of
10 example, it may be possible to yield similar or even substantially higher transformation frequencies.

- Without wishing to be bound by theory, we also believe that the degree of selectivity may differ among different varieties/cultivars/subspecies of a plant species. Also, other
15 physical and/or chemical and/or biological parameters affecting the effect of galactose may be expected to have an impact on the number of selectable transgenic shoots, possibly increasing the transformation frequencies significantly.

EXAMPLE 5 - TRANSGENIC RAPE SEED

- 20 Hypocotyl explants of oil seed rape were prepared using a protocol modified according to Radke *et al.* (1988. Theor. Appl. Genet. 78: 161-168). Hypocotyls of 5 d old seedlings were cut into 1 cm segments and inoculated with a suspension of *Agrobacterium tumefaciens* strain EHA101 ($OD_{660}=0.1-0.5$, prepared according to
25 standard procedures). After inoculation, the explants were dried carefully on filterpaper before co-cultivation.

Selection of transgenic oil seed rape on galactose containing media

After co-culture, the explants were transferred to selection medium consisting of modified MS-medium (Murashige and Skoog 1962. *Physiol. Plant.* 15: 473-497) supplemented with 500 mg/l carbenicillin and various concentrations of sucrose and galactose. After 7-8 weeks, shoots were harvested and transgenicity was evaluated by subjecting the harvested shoots to the assay for β -glucuronidase activity (Jefferson *et al.* 1987. *Plant Mol. Biol. Rep.* 5: 387-405), a marker enzyme encoded by a gene placed adjacent the selective gene on the T-DNA of the transformation vector (Fig. 3). The number of harvested shoots which were more than 50% β -glucuronidase-positive is indicated in the table below.

Galactose concentration (g/l)	Sucrose concentration (g/l)	No. of transgenic shoots	Transformation frequency (%)
0	20	0	0
2.5	10	4	1.0

The data show that it is possible to select transgenic shoots of oil seed rape, transformed with a gene encoding an enzyme that enables or enhances the ability of the transformed shoots to reduce the toxicity of galactose.

Without wishing to be bound by theory, we also believe that the degree of selectivity may differ due to the choice of the selection media - which may contain galactose and/or metabolic precursors and/or metabolic derivatives and/or derivatives of galactose, alone or in combination with other carbohydrate/energy source(s) such as sucrose, glucose, fructose, maltose which may or may not affect the effect of galactose. By way of example, it may be possible to yield similar or even substantially higher transformation frequencies.

Without wishing to be bound by theory, we also believe that the degree of selectivity may differ among different varieties/cultivars/subspecies of a plant species. Also, other physical and/or chemical and/or biological parameters affecting the effect of galactose may be expected to have an impact on the number of selectable transgenic shoots,
5 possibly increasing the transformation frequencies significantly.

SUMMATION

In summation, therefore, the present invention relates to a selection method for selecting
10 from a population of cells one or more selectable genetically transformed cells. The population of cells comprises selectable genetically transformed cells and possible non-transformed cells. Each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product. Each of the selectable genetically transformed cells optionally comprises an optional second
15 expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product. In the method a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the
20 optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product. The component can be present in an amount that is toxic to the non-transformed cells. The selection method comprises the step of introducing the population of cells to a medium, wherein the medium comprises the component and/or a derivative thereof and/or a precursor thereof
25 and in an amount such that the component is or will be in an amount that is utilisable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells. In the method, each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative
30 thereof or a precursor thereof. In the method, the component and/or the derivative

thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

The present invention also relates to a number of other aspects which also contain some
5 or all of the same distinguishing technical features - such as: a composition comprising a
population of cells comprising selectable genetically transformed cells and possible non-
transformed cells, and a medium; a population of cells comprising selectable genetically
transformed cells and possible non-transformed cells; a selectable genetically
transformed cell; an organism comprising a selectable genetically transformed cell; a
10 construct for genetically transforming a non-transformed cell to produce a selectable
genetically transformed cell.

In one aspect of the present invention, the component is present in the medium.

15 Hence, this aspect of the present invention can be expressed as:

a selection method for selecting from a population of cells one or more selectable
genetically transformed cells,

wherein the population of cells comprises selectable genetically transformed cells
and possible non-transformed cells;

20 wherein each of the selectable genetically transformed cells comprises a
first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises
an optional second expressable nucleotide sequence encoding a second expression
product and/or an optional third expressable nucleotide sequence encoding a third
25 expression product;

wherein a component is utilisable by the selectable genetically transformed cells
by action of the first expressable nucleotide sequence or the first expression product and
optionally by action of the optional second expressable nucleotide sequence or the
optional second expression product and/or by action of the optional third expressable
30 nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

the method comprising the step of introducing the population of cells to a medium,

wherein the medium comprises the component and optionally a derivative thereof
5 and/or a precursor thereof and in an amount such that the component is or will be in an amount that is utilisable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an
10 enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

15 Likewise, this aspect of the present invention also relates to a number of other aspects which also contain some or all of the same distinguishing technical features - such as: a composition comprising a population of cells comprising selectable genetically transformed cells and possible non-transformed cells, and a medium; a population of cells comprising selectable genetically transformed cells and possible non-transformed
20 cells; a selectable genetically transformed cell; an organism comprising a selectable genetically transformed cell; a construct for genetically transforming a non-transformed cell to produce a selectable genetically transformed cell.

For some applications the component is detoxifiable by the selectable genetically
25 transformed cells.

One aspect of this aspect of the present invention can be expressed as:

a selection method for selecting from a population of cells one or more selectable genetically transformed cells,

30 wherein the population of cells comprises selectable genetically transformed cells and possible non-transformed cells;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component or a derivative thereof or a precursor thereof when present in a detoxifiable amount in a medium is detoxifiable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component or the derivative thereof or the precursor thereof can be present in an amount in a medium that is toxic to the non-transformed cells; the method comprising the step of introducing the population of cells to a medium,

wherein the medium comprises the component and/or the derivative thereof and/or the precursor therefor and in an amount that is detoxifiable by the transformed cells but in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

25

Likewise, this aspect of the present invention also relates to a number of other aspects which also contain some or all of the same distinguishing technical features - such as: a composition comprising a population of cells comprising selectable genetically transformed cells and possible non-transformed cells, and a medium; a population of cells comprising selectable genetically transformed cells and possible non-transformed cells; a selectable genetically transformed cell; an organism comprising a selectable

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genetically transformed cell; a construct for genetically transforming a non-transformed cell to produce a selectable genetically transformed cell.

In one preferred aspect, the component is prepared *in situ* in the cells by processing a precursor therefor or a derivative thereof that was present in a medium in which the cells are or were present.

Hence, this preferred aspect can be expressed as:

a selection method for selecting from a population of cells one or more selectable genetically transformed cells,

wherein the population of cells comprises selectable genetically transformed cells and possible non-transformed cells;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component is utilisable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

the method comprising the step of introducing the population of cells to a medium,

wherein the medium comprises a derivative of the component and/or a precursor of the component and in an amount such that the component is or will be in an amount that is utilisable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

5 wherein the component and/or the derivative thereof and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

Likewise, this preferred aspect of the present invention also relates to a number of other aspects which also contain some or all of the same distinguishing technical features -
10 such as: a composition comprising a population of cells comprising selectable genetically transformed cells and possible non-transformed cells, and a medium; a population of cells comprising selectable genetically transformed cells and possible non-transformed cells; a selectable genetically transformed cell; an organism comprising a selectable genetically transformed cell; a construct for genetically transforming a non-transformed
15 cell to produce a selectable genetically transformed cell.

All publications mentioned in the above specification are herein incorporated by reference. Various modifications and variations of the described methods and system of the invention will be apparent to those skilled in the art without departing from the scope
20 and spirit of the invention. Although the invention has been described in connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes for carrying out the invention which are obvious to those skilled in molecular biology or related fields are intended to be within the scope of
25 the following claims.

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5 CLAIMS

1. A selection method for selecting from a population of cells one or more selectable genetically transformed cells,

wherein the population of cells comprises selectable genetically transformed
10 cells and possible non-transformed cells;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second
15 expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component is detoxifiable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide
20 sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

the method comprising the step of introducing the population of cells to a medium,

25 wherein the medium comprises the component and/or a precursor thereof and in an amount such that the component is or will be in an amount that is detoxifiable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second
30 expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the precursor thereof is galactose or a precursor thereof.

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2. A composition comprising a population of cells comprising selectable genetically transformed cells and possible non-transformed cells; and a medium;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component is detoxifiable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

wherein the medium comprises the component and/or a precursor thereof and in an amount such that the component is or will be in an amount that is detoxifiable by the transformed cells but wherein the component is or will be in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

5 3. A population of cells comprising selectable genetically transformed cells and possible non-transformed cells;

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells
10 comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

wherein a component is detoxifiable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide
15 sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

20 wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the precursor thereof is galactose or a precursor
25 thereof.

4. A selectable genetically transformed cell comprising a first expressable nucleotide sequence encoding a first expression product; and optionally a second expressable nucleotide sequence encoding a second expression product and/or a third
30 expressable nucleotide sequence encoding a third expression product;

wherein a component is detoxifiable by the selectable genetically transformed cells by action of the the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional
35 third expressable nucleotide sequence or the optional third expression product;

5 wherein the component can be present in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

10 wherein the component and/or the precursor thereof is galactose or a precursor thereof.

5. An organism comprising a selectable genetically transformed cell according to claim 4.

6. A construct for genetically transforming a non-transformed cell to produce a selectable genetically transformed cell;

the construct comprising a first expressable nucleotide sequence encoding a first expression product; and optionally a second expressable nucleotide sequence encoding a second expression product and/or a third expressable nucleotide sequence encoding a third expression product;

wherein a component is detoxifiable by the selectable genetically transformed cells by action of the first expressable nucleotide sequence or the first expression product and optionally by action of the optional second expressable nucleotide sequence or the optional second expression product and/or by action of the optional third expressable nucleotide sequence or the optional third expression product;

wherein the component can be present in an amount that is toxic to the non-transformed cells;

wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof; and

wherein the component and/or the precursor thereof is galactose or a precursor thereof.

- 5 7. A vector comprising the construct according to claim 6.
8. A plasmid comprising the construct according to claim 6.
9. A kit comprising a construct according to claim 6 or a vector according to
10 claim 7 or a plasmid according to claim 8 for genetically transforming a non-
transformed cell to produce a selectable genetically transformed cell; and a medium.
10. The invention according to any one of the preceding claims wherein the
selectable genetically transformed cell/cells is/are *in vitro* within a culture.
11. The invention according to any one of the preceding claims wherein the
selectable genetically transformed cell/cells is/are *in vivo* within an organism.
12. The invention according to any one of the preceding claims wherein the
selectable genetically transformed cell/cells is/are selectable genetically transformed
plant cell/cells.
13. The invention according to any one of the preceding claims wherein an
additional nucleotide sequence is present and wherein the additional nucleotide
sequence codes for a nucleotide sequence of interest.
14. The invention according to any one of the preceding claims wherein the
component is present in the medium.
15. The invention according to any one of claims 1 to 13 wherein the component
is prepared *in situ* in the cell from a precursor that was present in the medium.
16. A plant prepared from or comprising the invention according to any one of the
preceding claims, preferably wherein the plant is capable of providing a feed,
foodstuff to humans or animals.

5 17. A plant according to claim 14 wherein the plant is any one of rape seed, potato or maize.

18. A plant comprising one or more heterologous enzymes, wherein the heterologous enzymes are any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10) and/or UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12).

15 19. Use of any one or more of galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) as a selection means for selecting a genetically transformed plant cell over a non-transformed plant cell.

20 20. Use of any one or more of a nucleotide sequence coding for galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12), UDP-galactose epimerase (EC 5.1.3.2) for providing a selection means for selecting a genetically transformed plant cell over a non-transformed plant cell.

25 21. Use of a prokaryote comprising a nucleotide sequence coding for galactokinase (EC 2.7.1.6), UTP-dependent pyrophosphorylase (EC 2.7.7.10), UDP-glucose-dependent uridylyltransferase (EC 2.7.7.12) and/or UDP-galactose epimerase (EC 5.1.3.2); wherein the nucleotide sequence comprises at least one intron which inactivates the nucleotide sequence or the expression product thereof in the prokaryote, for providing a selection means for selecting a genetically transformed cell over a non-transformed cell.

22. Use of any one or more of galactose, galactose-1-phosphate, UDP-galactose, or a derivative thereof, as a selection means for selecting a genetically transformed plant cell over a non-transformed plant cell.

35

23. A selection system that uses a component or a precursor therefor or a derivative thereof for selecting at least one genetically transformed cell from a

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5 population of cells, wherein the at least one genetically transformed cell is transformed with a nucleotide sequence which encodes an expression product capable of utilising the component but wherein the component toxic to non-transformed cells, wherein the component and/or the precursor thereof is galactose or a derivative thereof or a precursor thereof.

10

24. A selection method for selecting from a population of cells one or more selectable genetically transformed cells,

wherein the population of cells comprises selectable genetically transformed cells and possible non-transformed cells;

15

wherein each of the selectable genetically transformed cells comprises a first expressable nucleotide sequence encoding a first expression product;

optionally wherein each of the selectable genetically transformed cells comprises an optional second expressable nucleotide sequence encoding a second expression product and/or an optional third expressable nucleotide sequence encoding a third expression product;

20

the method comprising the step of preparing the population of cells by transforming some or all of the cells in an initial population of cells containing non-transformed cells with a heterologous nucleotide sequence so as to form the population of cells containing one or more selectable genetically transformed cells;

25

and

selecting at least one of the selectable genetically transformed cells;

wherein the heterologous nucleotide sequence is any one or more of the first expressable nucleotide sequence, the optional second expressable nucleotide sequence or the optional third expressable nucleotide sequence;

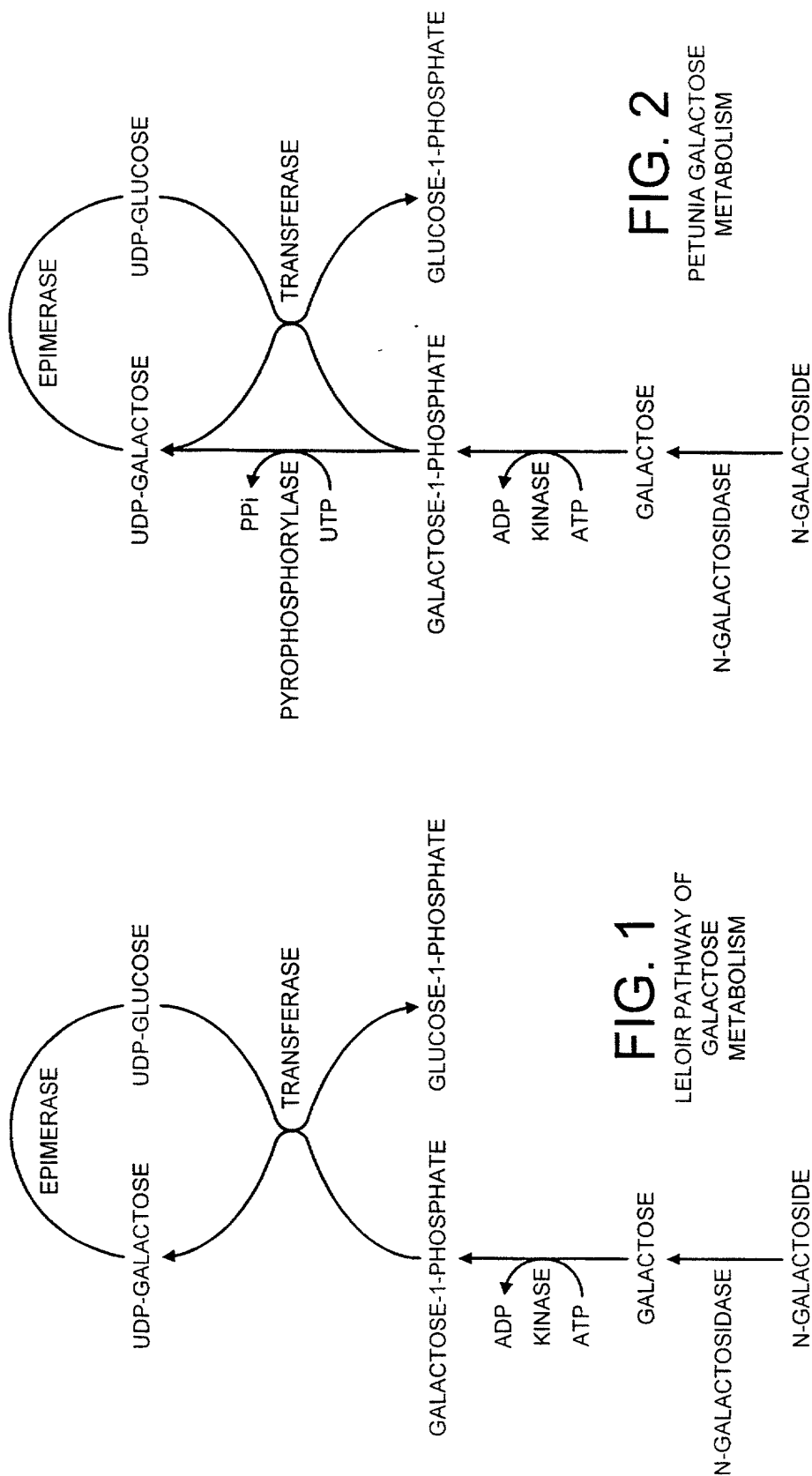
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- 5 wherein each of the first expression product and the optional second expression product and the optional third expression product is independently selected from an enzyme capable of metabolising galactose or a derivative thereof or a precursor thereof.

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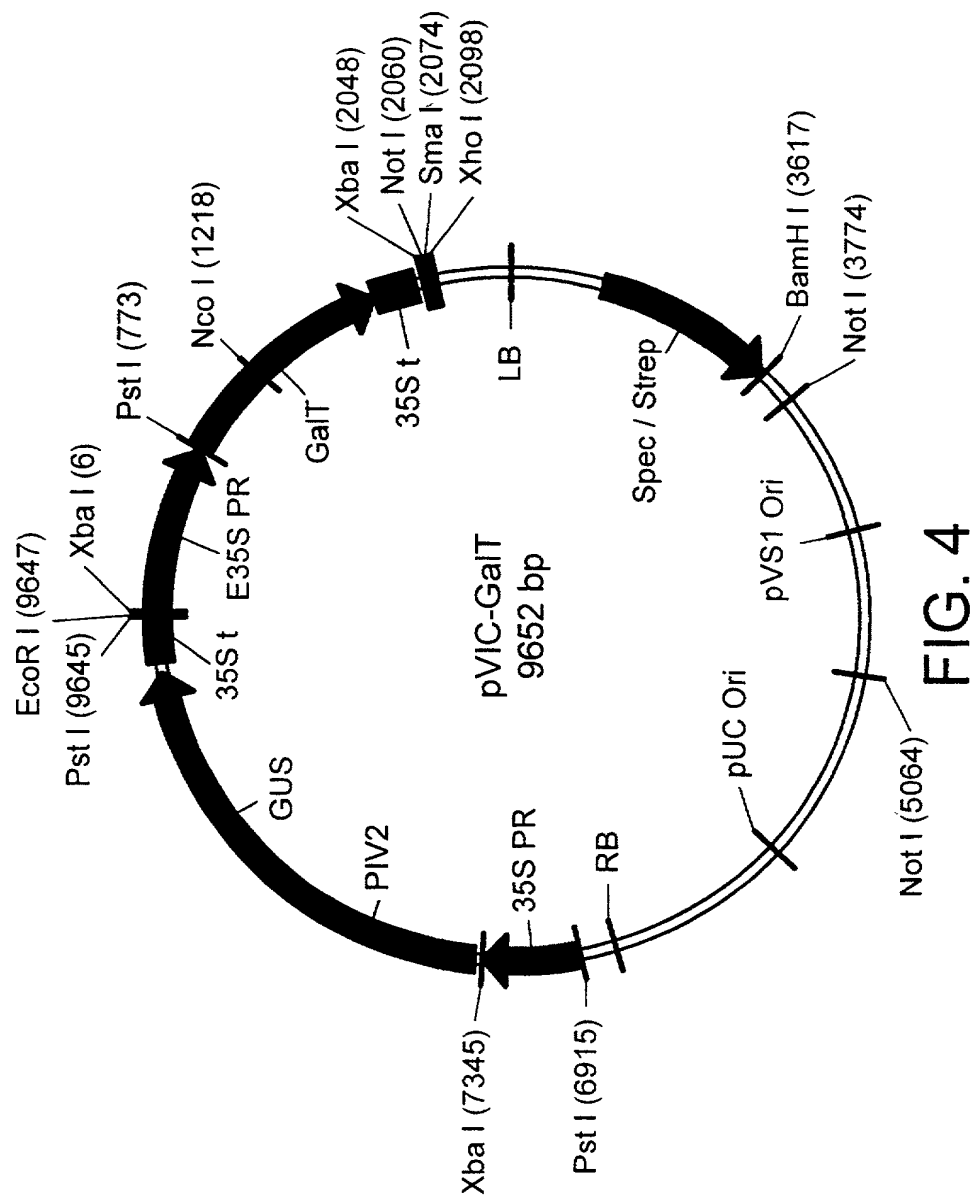
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 W P F E T L L L P K A H V L R I T D L T D A Q R S D L A L A L K K L 262

 GACCAGTCGTTATGACAAACCTCTTCCAGTGCTCTTCCCTACTCTATGGGCTGGCACGGCGGCCATTTAATGGCGAAGAGAATCAACACTGGCAGCTG 900
 T S R Y D N L F Q C S F P Y S M G W H G A P F N G E E N Q H W Q L 295

 CACGGGCACTTTTATCCGCTCTGCTGCGTCCGCCACGTACGTAATTTATGGTTGTTATGAATGCTGGCAGAGACCCAGCAGACCTGACCCGAG 1000
 H A H F Y P P L L R S A T V R K F M V G Y E M L A E T Q R D L T A 328

 AACAGGACGAGAGCGTTTGGCGCAGTCAGCGATATCCATTTTCGGAATCCGGAGTGAAGTCGACT 1069
 E Q A A E R L R A V S D I H F R E S G V - Sali 348

FIG. 3



United States Patent Application

COMBINED DECLARATION AND POWER OF ATTORNEY

As a below named inventor I hereby declare that: my residence, post office address and citizenship are as stated below next to my name; that

I verily believe I am the original, first and sole inventor (if only one name is listed below) or a joint inventor (if plural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: **SELECTION METHOD**

The specification of which

- a. ☒ is attached hereto
b. ☐ was filed on _____ as application serial no. _____ and was amended on _____ (if applicable) (in the case of a PCT-filed application) described and claimed in international no. PCT/IB99/01465 filed August 11, 1999 and as amended on _____ (if any), which I have reviewed and for which I solicit a United States patent.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I hereby claim foreign priority benefits under Title 35, United States Code, § 119/365 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on the basis of which priority is claimed:

- a. ☐ no such applications have been filed.
b. ☒ such applications have been filed as follows:

FOREIGN APPLICATION(S), IF ANY, CLAIMING PRIORITY UNDER 35 USC § 119			
COUNTRY	APPLICATION NUMBER	DATE OF FILING (day, month, year)	DATE OF ISSUE (day, month, year)
Great Britain	9817465.9	August 11, 1998	
ALL FOREIGN APPLICATION(S), IF ANY, FILED BEFORE THE PRIORITY APPLICATION(S)			
COUNTRY	APPLICATION NUMBER	DATE OF FILING (day, month, year)	DATE OF ISSUE (day, month, year)

I hereby claim the benefit under Title 35, United States Code, § 120/365 of any United States and PCT international application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application.

U.S. APPLICATION NUMBER	DATE OF FILING (day, month, year)	STATUS (patented, pending, abandoned)

I hereby claim the benefit under Title 35, United States Code § 119(e) of any United States provisional application(s) listed below:

U.S. PROVISIONAL APPLICATION NUMBER	DATE OF FILING (Day, Month, Year)

§ 1.56 Duty to disclose information material to patentability.

(d) A patent by its very nature is affected with a public interest. The public interest is best served, and the most effective patent examination occurs when, at the time an application is being examined, the Office is aware of and evaluates the teachings of all information material to patentability. Each individual associated with the filing and prosecution of a patent application has a duty of candor and good faith in dealing with the Office, which includes a duty to disclose to the Office all information known to that individual to be material to patentability as defined in this section. The duty to disclose information exists with respect to each pending claim until the claim is canceled or withdrawn from consideration, or the application becomes abandoned. Information material to the patentability of a claim that is canceled or withdrawn from consideration need not be submitted if the information is not material to the patentability of any claim remaining under consideration in the application. There is no duty to submit information which is not material to the patentability of any existing claim. The duty to disclose all information known to be material to patentability is deemed to be satisfied if all information known to be material to patentability of any claim issued in a patent was cited by the Office or submitted to the Office in the manner prescribed by §§ 1.97(b)-(d) and 1.98. However, no patent will be granted on an application in connection with which fraud on the Office was practiced or attempted or the duty of disclosure was violated through bad faith or intentional misconduct. The Office encourages applicants to carefully examine:

- (1) prior art cited in search reports of a foreign patent office in a counterpart application, and
- (2) the closest information over which individuals associated with the filing or prosecution of a patent application believe any pending claim patentably defines, to make sure that any material information contained therein is disclosed to the Office

(b) Under this section, information is material to patentability when it is not cumulative to information already of record or being made of record in the application, and

- (1) It establishes, by itself or in combination with other information, a prima facie case of unpatentability of a claim;

or

- (2) It refutes, or is inconsistent with, a position the applicant takes in:
 - (i) Opposing an argument of unpatentability relied on by the Office; or
 - (ii) Asserting an argument of patentability.

A prima facie case of unpatentability is established when the information compels a conclusion that a claim is unpatentable under the preponderance of evidence, burden-of-proof standard, giving each term in the claim its broadest reasonable construction consistent with the specification, and before any consideration is given to evidence which may be submitted in an attempt to establish a contrary conclusion of patentability.

- (c) Individuals associated with the filing or prosecution of a patent application within the meaning of this section are:
 - (1) Each inventor named in the application;
 - (2) Each attorney or agent who prepares or prosecutes the application; and
 - (3) Every other person who is substantively involved in the preparation or prosecution of the application and who is associated with the inventor, with the assignee or with anyone to whom there is an obligation to assign the application.
- (d) Individuals other than the attorney, agent or inventor may comply with this section by disclosing information to the attorney, agent, or inventor.
- (e) In any continuation-in-part application, the duty under this section includes the duty to disclose to the Office all information known to the person to be material to patentability, as defined in paragraph (b) of this section, which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby appoint the following attorney(s) and/or patent agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected herewith:

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I hereby authorize them to act and rely on instructions from and communicate directly with the person/assignee/attorney/firm/ organization who/which first sends/sent this case to them and by whom/which I hereby declare that I have consented after full disclosure to be represented unless/until I instruct Merchant & Gould P.C. to the contrary.

Please direct all correspondence in this case to Merchant & Gould P.C. at the address indicated below:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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